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(54) Title: METHODS AND COMPOSITIONS FOR INHIBITING CD14 MEDIATED CELL ACTIVATION		
(57) Abstract <p>This invention provides hybridoma cell lines producing monoclonal antibodies which inhibit CD14 mediated cell activation. Monoclonal antibodies produced by these cell lines also are provided. The antibodies are useful for the detection of the presence of cell surface and soluble CD14 in a sample. Chimeric and CDR grafted antibodies generated from the above monoclonal antibodies are further provided. Pharmaceutical compositions containing the above biological compositions are provided. These are useful to treat and prevent disorders associated with CD14 mediated cell activation, such as spesis.</p>		

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**METHODS AND COMPOSITIONS FOR
INHIBITING CD14 MEDIATED CELL ACTIVATION**

This application is a continuation of application Serial No. 08/070,160, filed on May 28, 1993.

5 BACKGROUND OF THE INVENTION

1. Field of the Invention

The present invention relates to methods and compositions for inhibiting CD14 mediated cell activation. More particularly, the present invention relates to molecules that bind the CD14 monocyte antigen at a site which inhibits CD14 mediated cell activation.

2. Description of the Related Art

The correct functioning of a cell depends partly on its ability to communicate with its environment; external stimuli often interact with membrane receptors which, in turn, induce second messengers that ultimately affect transcription factors. The latter then activate or repress the expression of certain genes resulting in a specific pattern of proteins in a given cell.

The transcription factor NF- κ B (nuclear factor- κ B) is induced by a variety of stimuli to contact its DNA-binding motif and regulate a set of genes encoding immunoreceptors, cytokines, and viral proteins. Included among the various factors which can activate NF- κ B is lipopolysaccharide (LPS). LPS, in turn, is intimately involved in the induction of the sepsis syndrome, including septic shock, systemic inflammatory response syndrome, and multiorgan failure.

Sepsis is a morbid condition induced by a toxin, the introduction or accumulation of which is most commonly caused by infection or trauma. The initial symptoms of sepsis typically include chills, profuse sweat, irregularly

remittent fever, prostration and the like, followed by persistent fever, hypotension leading to shock, neutropenia, leukopenia, disseminated intravascular coagulation, acute respiratory distress syndrome and multiple organ failure.

5 Lipopolysaccharide, or endotoxin, is a toxic component found in the outer membrane of all gram-negative microorganisms (e.g., *Escherichia coli*, *Klebsiella pneumonia*, *Pseudomonas aeruginosa*). It has been determined that LPS is a potent and pleiotropic stimulus for immune cells, both *in vitro* and *in vivo* (Morrison, D.C. & J.L. Ryan, *Annu. Rev. Med.*, 38:417, 1987; Bone, R.C.,
10 *Ann. Intern. Med.*, 115:457, 1991). Compelling evidence supports the toxic role of LPS in that all of the pathophysiological effects noted in humans during gram-negative sepsis can be completely duplicated with purified LPS. The mechanism by which this toxic component activates responsive cells is complex and not fully understood. The host response to gram-negative
15 bacterial infection is dependent upon effector cell recognition of these bacteria and/or LPS and involves serum proteins and cell membrane receptors. While the clearance of bacteria and LPS is via endocytosis and phagocytosis by reticuloendothelial cells, concomitant activation of the host immune response by LPS results in secretion of cytokines by activated macrophages which can
20 trigger the exaggerated host responses that occur during gram-negative bacterial infection.

The discovery by Tobias, *et al.* (*J. Exp. Med.*, 164:777, 1986) of a serum protein, identified as LPS binding protein (LBP), that exhibits high affinity binding to LPS ($K_d \approx 10^{-9} \text{ M}^{-1}$), helped to define the fate of LPS once released
25 *in vivo*. It was demonstrated that this novel protein, with a molecular weight of 60kD, which is synthesized in the liver is an acute phase serum protein reaching levels of 200 μ g/ml in humans. The formation of high affinity LPS/LBP complexes is followed by recognition by macrophages with subsequent release

of $\text{TNF-}\alpha$ and other macrophage secretory products (Schumann, R.R., *et al.*, *Science*, 249:1429, 1990). Additional studies on the effects of LPS complexed with LPB led to the discovery of its specific receptor on the surface of monocytes and macrophages; CD14 (Wright, S.D., *et al.*, *Science*, 249:1431, 1990). Further analysis with mAbs specific for CD14 revealed that the domain to which one anti-CD14 mAb (3C10; VanVoorhis, W.C., *et al.*, *J. Exp. Med.*, 158:126, 1983) bound was part of, or in close proximity to, the LPS/LBP binding site on CD14. Monoclonal antibody 3C10, by nature of its ability to block LPS/LBP binding to CD14, was capable of inhibiting $\text{TNF-}\alpha$ release in a human whole blood assay, after stimulation with LPS. It is suggested by this discovery that the blocking of a single protein determinant (the ligand binding site on CD14) is sufficient, even in the presence of all other cells, proteins and factors contained in human whole blood, to inhibit $\text{TNF-}\alpha$ release (known to be a key mediator in septic shock) and other macrophage secretory products in response to LPS.

In spite of the advances which have been made in understanding the nature of CD14 mediated cell activation disorders, such as sepsis, considerable need remains for compositions which can be used to inhibit such activation and to diagnose these disorders. The present invention provides such compositions.

SUMMARY OF THE INVENTION

This invention provides hybridoma cell lines producing monoclonal antibodies, the monoclonal antibodies being capable of inhibiting CD14 mediated cell activation. Monoclonal antibodies produced by these cell lines also are
5 provided. These monoclonal antibodies are broadly useful in inhibiting NF- κ B activation by a ligand which binds to CD14 and would otherwise be capable of inducing NF- κ B activation. Biologically active fragments of the monoclonal antibodies are provided. The antibodies and fragments are useful for the
10 detection of the presence of cell surface-associated and soluble CD14 in a sample. Chimeric and CDR grafted antibodies generated from the above monoclonal antibodies are further provided.

Pharmaceutical compositions containing the above biological compositions are provided. These are useful to treat and prevent LPS-associated disorders, such as sepsis.

BRIEF DESCRIPTION OF THE FIGURES

Figure 1 is the nucleic acid sequence for the human soluble CD14 receptor (SEQ ID NO:9).

5 Figure 2 is the nucleic acid and amino acid sequence of the 28C5 heavy chain (SEQ ID NOS:1 and 2, respectively).

Figure 3 is the nucleic acid and amino acid sequence of the 28C5 light chain (SEQ ID NOS:3 and 4, respectively).

Figure 4 is the nucleic acid and amino acid sequence of the 18E12 heavy chain (SEQ ID NOS:5 and 6, respectively).

10 Figure 5 is the nucleic acid and amino acid sequence of the 18E12 light chain (SEQ ID NOS:7 and 8, respectively).

Figure 6 is FACS analysis of control THP-1 cells only.

Figure 7 is FACS analysis of control THP-1 cells and FITC conjugate only.

Figure 8 is FACS analysis of positive control THP-1 and MY4 antibody.

15 Figure 9 is FACS analysis of 28C5 antibody.

Figure 10 is FACS analysis of 18E12 antibody.

Figure 11 shows the titers of various mAb to sCD14.

Figure 12 shows the results of a competition assay between mAb 3C10 and a panel of anti-CD14 mAbs.

Figure 13 shows the blocking of LPS/LBP binding to CD14 by anti-CD14 mAbs.

5 Figure 14 shows the results of an evaluation of the ability of anti-CD14 mAbs to block cytokine release in HL-60 cells.

Figure 15 shows effect of anti-CD14 mAbs to inhibit LPS binding to cellular CD14.

Figure 16 shows effect of anti-CD14 mAbs on LPS-dependent, CD14-mediated activation of cells.

10 Figure 17 shows the mean arterial pressure of monkeys challenged with LPS and treated with 18E12 (•), 28C5 (■) or IgG1 (x).

Figure 18 shows pre and post human IFN- γ treatment CD14 levels and LBP levels in a monkey.

15 Figure 19 shows the lavage/plasma ration of BSA in monkeys treated with 18E12, 28C5 or IgG1.

Figure 20 shows the antibody half life of 18E12, 28C5 and IgG1 in monkeys.

Figure 21 shows CD14 levels in monkeys treated with antibody (18E12, 28C5 and IgG1) alone (top) or challenged with LPS after antibody treatment (bottom).

Figure 22 shows LBP levels in monkeys treated with antibody (18E12, 28C5 and IgG1) alone (top) or challenged with LPS after antibody treatment (bottom).

5 Figure 23 shows ALT/GPT levels in monkeys treated with antibody (18E12, 28C5 and IgG1) alone (top) or challenged with LPS after antibody treatment (bottom).

Figure 24 shows E-selectin levels in monkeys treated with antibody (18E12, 28C5 and IgG1) alone (top) or challenged with LPS after antibody treatment (bottom).

10 Figure 25 shows TNF levels in monkeys treated with antibody (18E12, 28C5 and IgG1) and challenged with LPS (top) and IL-1 levels in monkeys treated with antibody (18E12, 28C5 and IgG1) and challenged with LPS after antibody treatment (bottom).

15 Figure 26 shows IL-6 levels in monkeys treated with antibody (18E12, 28C5 and IgG1) and challenged with LPS (top) and IL-8 levels in monkeys treated with antibody (18E12, 28C5 and IgG1) and challenged with LPS after antibody treatment (bottom).

Figure 27 shows inhibition of TNF release in human whole blood stimulated with LPS by treatment with 18E12 (•), 28C5 (▲) or 23G4 (■).

20 Figure 28 shows the effect of anti-CD14 antibodies on LPS-induced TNF secretion in baboon blood. (23G4 (•), 28C5 (▽), and 18E12 (▼).

Figure 29 shows the amino acid sequence of the light chains of monoclonal antibodies 3C10, 28C5, 23G4 and 18E12.

Figure 30 shows the amino acid sequence of the heavy chains of monoclonal antibodies 3C10, 28C5, and 18E12.

DETAILED DESCRIPTION OF THE INVENTION

A full length polypeptide for a human soluble CD14 ("sCD14") is provided by the disclosure. As used herein, "CD14" means the cell surface receptor that has been identified as the binding site for LPS when the LPS is present as an LPS:LBP complex. The CD14 cell surface receptor is a glycerophosphatidylinositol (GPI)-linked protein present on the surface of mature monocytes, neutrophils, and macrophages. Native CD14 also is spontaneously released from the surface of mature monocytes and macrophages in a soluble form. Native sCD14 lacks the GPI anchor and is present in serum. The biological origin and function of sCD14 have not yet been fully defined (Bazil, *Europ. J. Immunol.*, 16:1583-1589, 1986).

As used herein, "soluble" is defined as not associated in the cell surface. "Soluble CD14" is a non-cell-associated CD14 molecule further characterized as specifically binding LPS:LBP complexes and/or LPS alone. "Recombinant human sCD14" includes both a full-length amino acid soluble human CD14 protein encoded by the nucleic acid sequence in Figure 1 (SEQ ID NO:9) and its truncated version. For the purposes of identification only, the full-length protein is designated 523 and the truncated version is designated 847. This human sCD14 is useful as an immunogen for the generation of polyclonal and monoclonal antibodies and to detect the presence of LPS in a patient sample. When used as an immunogen, 523 provided advantages over prior art CD14 immunogens, e.g., 523 provided a higher number of CD14-specific positive clones, it eliminated the number of non-specific responses to other immunogenic proteins which would be present in whole cell extracts, and it decreased the number of screening attempts needed to obtain the antibodies of interest. This full length human sCD14 polypeptide has the nucleic acid sequence as shown in Figure 1, SEQ ID NO:9. Using this sequence, one of skill in the art can produce polypeptide of like sequence by chemical synthesis

or recombinantly. The truncated version ("847") has eight (8) amino acids from the carboxyl end of the sequence deleted.

Minor modifications of sCD14 primary amino acid sequence may result in proteins which have substantially equivalent function compared to the sCD14 protein described herein. Such modifications may be deliberate, as by site-directed mutagenesis, or may be spontaneous. All proteins produced by these
5 modifications are included herein as long as sCD14 function exists.

Modifications of sCD14 primary amino acid sequence also include conservative variations. The term "conservative variation" as used herein denotes the replacement of an amino acid residue by another, biologically similar residue.
10 Examples of conservative variations include the substitution of one hydrophobic residue such as isoleucine, valine, leucine or methionine for another, or the substitution of one polar residue for another, such as the substitution of arginine for lysine, glutamic for aspartic acids, or glutamine for asparagine, and
15 the like. The term "conservative variation" also includes the use of a substituted amino acid in place of an unsubstituted parent amino acid provided that antibodies raised to the substituted polypeptide also immunoreact with the unsubstituted polypeptide.

The invention provides a nucleic acid molecule encoding the human soluble CD14 polypeptide as shown in Figure 1 (SEQ ID NO:9). The invention also encompasses nucleic acids molecules which differ from that of the nucleic acid molecule shown in Figure 1, but which produce the same phenotypic or immunogenic effect when the nucleic acid molecule is expressed. This invention encompasses nucleic acid molecules characterized by changes in
20 non-coding regions that do not alter the phenotype of the polypeptide produced therefrom when compared to the nucleic acid molecule described hereinabove. Therefore, it is understood that all polynucleotides encoding all
25

or a portion of sCD14 are also included herein, so long as they exhibit a function of sCD14, such as the ability to induce or bind antibody. Such polynucleotides include both naturally occurring and intentionally manipulated, for example, mutagenized polynucleotides. These polynucleotides include DNA and RNA sequences which encode the protein.

This invention further encompasses nucleic acid molecules which hybridize to the nucleic acid molecule of the subject invention. As used herein, the term "nucleic acid" encompasses RNA as well as single and double stranded DNA and cDNA.

Using the sequence provided in Figure 1 and methods well known to those of skill in the art (as exemplified in Sambrook, *et al.*, *Molecular Cloning: A Laboratory Manual*, Cold Spring Harbor Laboratory, New York (1989), incorporated throughout by reference), human sCD14 can be recombinantly produced and isolated. Expression vectors containing this sequence, as well as host cells containing the vectors, are also provided by this invention. As used herein, the term "vector" or "expression vector" refers to sequences of heterologous nucleic acids which are capable of being expressed in selected host cells through operational association with other sequences capable of effecting their expression, such as promoter and enhancer sequences. For the purpose of illustration only, these expression vectors can be bacterial plasmids, bacterial phages, animal viruses, baculoviruses or cosmids. Procaryotic host cells such as *E. coli* can be used for recombinantly producing these polypeptides when the vector is a bacterial plasmid or a bacterial phage. Eucaryotic host cells can be, but are not limited to mammalian host cells, e.g., Chinese Hamster Ovary Cells (CHO) or insect cells for baculoviral expression.

A method of recombinantly producing the human sCD14 is provided by this invention. This method requires growing the host cells described above under suitable conditions such that the sCD14 nucleic acid molecule is transcribed and translated. Upon expression, the recombinant sCD14 can be isolated from
5 the cell culture by use of an affinity column composed of commercially available CD14 monoclonal antibody.

This invention also provides polyclonal antibodies and monoclonal antibodies, specifically reactive with cell surface CD14 receptor and soluble CD14. The antibodies of the invention inhibit CD14 mediated cell activation by a ligand
10 otherwise capable of binding to the CD14 receptor and activating the cell, for example, to induce NK- κ B activation or produce and release a cytokine. Monoclonal antibodies provided herein are capable of inhibiting CD14 mediated cell activation by the ligand even when the ligand has bound to CD14. The
15 monoclonal antibodies may allow at least about 50% ligand binding to occur between the ligand and CD14, although these antibodies can allow at least about 80% binding of ligand to CD14 to occur and still be capable of inhibiting CD14 mediated cell activation.

As used herein, a "antibody or polyclonal antibody" means a protein that is produced in response to immunization with an antigen or through recombinant
20 cloning techniques. The term "monoclonal antibody" means an immunoglobulin derived from a single clone of cells. All monoclonal antibodies derived from the clone are chemically and structurally identical, and specific for a single antigenic determinant.

Laboratory methods for producing polyclonal antibodies and monoclonal
25 antibodies are known in the art (see, Harlow and Lane, *Antibodies: A Laboratory Manual*, Cold Spring Harbor Laboratory, New York (1988), incorporated herein by reference). The monoclonal antibodies of this invention

can be biologically produced by introducing full length human recombinant polypeptide for soluble CD14 into an animal, e.g., a mouse or a rabbit. The antibody producing cells in the animal are isolated and fused with myeloma cells or heteromyeloma cells to produce hybrid cells or hybridomas.

5 Accordingly, the hybridoma cells producing the monoclonal antibodies of this invention also are provided. Monoclonal antibodies produced in this manner include, but are not limited to the monoclonal antibodies designated 18E12, 28C5, 23G4, 5G3, 4F2, 13A7, 10B7, and 26F3. The hybridoma cell lines 18E12, 28C5 and 23G4 have been deposited with the American Type Culture

10 Collection (ATCC) 12301 Parklawn Drive, Rockville, Maryland 20852, U.S.A., under the provisions of the Budapest Treaty on the International Deposit of Microorganisms for the Purposes of Patent Procedure. 18E12 and 28C5 were deposited on May 27, 1993 and were accorded ATCC accession numbers HB11363 and HB11364, respectively. 23G4 was deposited on May 25, 1994

15 and was accorded ATCC accession number X. These deposits were made under the provisions of the Budapest Treaty on the International Recognition of the Deposit of Microorganisms for the Purpose of Patent Procedure and the Regulations thereunder (Budapest Treaty). This assures maintenance of viable cultures for 30 years from the date of deposit. The organisms will be made

20 available by ATCC under the terms of the Budapest Treaty which assures permanent and unrestricted availability of the progeny of the culture to the public upon issuance of the pertinent U.S. patent or upon laying open to the public of any U.S. or foreign patent application, whichever comes first, and assures availability of the progeny to one determined by the U.S.

25 Commissioner of Patents and Trademarks to be entitled thereto according to 35 U.S.C. §122 and the Commissioner's rules pursuant thereto (including 37 C.F.R. §1.14 with particular reference to 886 OG 638).

5 The assignee of the present application has agreed that if the culture deposit should die or be lost or destroyed when cultivated under suitable conditions, it will be promptly replaced on notification with a viable specimen of the same culture. Availability of a deposited strain is not to be construed as a license to practice the invention in contravention of the rights granted under the authority of any government in accordance with its patent laws.

10 The foregoing written specification is considered to be sufficient to enable one skilled in the art to practice the invention. The present invention is not to be limited in scope by the cell lines deposited, since the deposited embodiment is intended as a single illustration of one aspect of the invention and any cell lines that are functionally equivalent are within the scope of this invention. The deposit of material does not constitute an admission that the written description herein contained is inadequate to enable the practice of any aspect of the invention, including the best mode thereof, nor is it to be construed as limiting
15 the scope of the claims to the specific illustration that it represents.

Thus, using the unique full length recombinant protein for soluble CD14 and the well known methods, one of skill in the art can produce and screen the hybridoma cells and antibodies of this invention.

20 The monoclonal antibodies of this invention can be characterized as being able to specifically react with cell surface CD14 receptor and soluble CD14 and inhibit CD14 mediated cell activation. Methods of determining binding specificities are outlined below. In one embodiment, the monoclonal antibodies are further characterized as having a binding affinity for CD14 greater than the affinity of anti-CD14 antibody 3C10 (available from the American Type Culture
25 Collection). One such monoclonal antibody is the monoclonal antibody designated 28C5. Scatchard analysis of 28C5 binding to dihydroxyvitamin D3 induced THPI cells gave an affinity of $3 \times 10^{-9} \text{M}^{-1}$. Monoclonal antibody 28C5

and 23G4, and antibodies of like specificity and affinity, are further characterized as being able to inhibit activation and also inhibit CD14 binding of the ligand which induces NF- κ B activation. In addition, all of the monoclonal antibodies of the invention can be characterized by their ability to inhibit cytokine release from CD14+ cells when such cells are contacted with the inducing ligand. As used herein, a cytokine shall include, but is not limited to TNF- α , IL-1, IL-6, and IL-8.

In an alternative embodiment, the monoclonal antibody 18E12 and monoclonal antibodies of like specificity are further characterized as having the ability to inhibit CD14 mediated cell activation, but do not significantly inhibit CD14 binding (i.e., these antibodies allow CD14 binding) with the ligand which is otherwise capable of inducing CD14 mediated cell activation. Monoclonal antibodies with the specificity of 18E12 will allow from at least about 50% to at least about 80% binding to occur between the ligand and CD14.

The preferred monoclonal antibodies described herein, 18E12 and 23G4, bind to both human and baboon CD14, whereas, 28C5 does not bind baboon CD14.

Although LBP is the predominant serum protein involved in presentation of LPS to CD14, other serum proteins may also bind to LPS under appropriate conditions and facilitate LPS-CD14 interactions (Wright, S.D., *et al.*, *J. Expt. Med.*, 176:719-727, 1992). Regardless of whether LBP or other proteins predominate under physiologic conditions the effects of the monoclonal antibodies 18E12, 23G4 or 28C5 are the same since these antibodies prevent the effects of LPS on NF- κ B or cytokine production in the presence plasma (or serum).

This invention also provides biological active fragments of the polyclonal and

monoclonal antibodies described above. These "antibody fragments" retain some ability to selectively bind with its antigen or receptor. Such antibody fragments can include, but are not limited to:

- 5 (1) Fab, the fragment which contains a monovalent antigen-binding fragment of an antibody molecule produced by digestion with the enzyme papain to yield an intact light chain and a portion of one heavy chain;
- 10 (2) Fab', the fragment of an antibody molecule obtained by treating with pepsin, followed by reduction, to yield an intact light chain and a portion of the heavy chain; two Fab' fragments are obtained per antibody molecule;
- (3) (Fab')₂, the fragment of the antibody that is obtained by treating with the enzyme pepsin without subsequent reduction; F(ab')₂ is a dimer of two Fab' fragments held together by two disulfide bonds;
- 15 (4) Fv, defined as a genetically engineered fragment containing the variable region of the light chain and the variable region of the heavy chain expressed as two chains; and
- 20 (5) Single chain antibody ("SCA"), defined as a genetically engineered molecule containing the variable region of the light chain, the variable region of the heavy chain, linked by a suitable polypeptide linker as a genetically fused single chain molecule.

Methods of making these fragments are known in the art, see for example, Harlow and Lane, *supra*.

Additional examples of "biologically active fragment" include antibody fragments specifically including the CDRs of the antibodies as defined below. These CDR regions are identified in Figures 2 through 5 and 29-30, (SEQ ID NOS:1 through 8 and SEQ ID NOS:22-24). CDRs of these antibodies are useful to generate CDR grafted antibodies as described below. Additional examples of "biologically active fragments" include fragments specifically including the framework regions of the antibodies also identified in Figures 2 through 5 and 29-30, (SEQ ID NOS:1 through 8 and SEQ ID NOS:22-24). The framework regions of the antibodies are useful as primers for PCR amplification of the CDRs.

Also encompassed by this invention are proteins or polypeptides that have been recombinantly produced, biochemically synthesized, chemically synthesized or chemically modified, that retain the ability to bind CD14 cell surface receptor and soluble CD14 as well as inhibit CD14 mediated cell activation by binding of activating ligand to CD14+ cells, of the corresponding native polyclonal or monoclonal antibody. The ability to bind with an antigen or receptor is determined by antigen-binding assays known in the art such as antibody capture assays (see, for example, Harlow and Lane, *supra*).

Any of the above described antibodies or biologically active antibody fragments can be used to generate CDR grafted and chimeric antibodies.

"CDR" or "complementarity determining region" or "hypervariable region" is defined as the amino acid sequences on the light and heavy chains of an antibody which form the three-dimensional loop structure that contributes to the formation of the antigen binding site.

As used herein, the term "CDR grafted" antibody refers to an antibody having an amino acid sequence in which at least parts of one or more CDR

sequences in the light and/or variable domain have been replaced by analogous parts of CDR sequences from an antibody having a different binding specificity for a given antigen or receptor.

5 As used herein, the terms "light chain variable region" and "heavy chain variable region" refer to the regions or domains at the N-terminal portion of the light and heavy chains respectively which have a varied primary amino acid sequence for each antibody. The variable region of the antibody consists of the amino terminal domain of the light and heavy chains as they fold together to form a three-dimensional binding site for an antibody.

10 The analogous CDR sequences are said to be "grafted" onto the substrate or recipient antibody. The "donor" antibody is the antibody providing the CDR sequence, and the antibody receiving the substituted sequences is the "substrate" antibody. One of skill in the art can readily produce these CDR grafted antibodies using the teachings provided herein in combination with
15 methods well known in the art (see Borrebaeck, C.A., *Antibody Engineering: A Practical Guide*, W.H. Freeman and Company, New York, 1992, incorporated throughout by reference).

This invention further provides chimeric antibodies of the above described antibodies or biologically active fragments. As used herein, the term "chimeric
20 antibody" refers to an antibody in which the variable regions of antibodies derived from one species are combined with the constant regions of antibodies derived from a different species. Chimeric antibodies are constructed by recombinant DNA technology, and are described in Shaw, *et al.*, *J. Immun.*, 138:4534 (1987), Sun, L.K., *et al.*, *Proc. Natl. Acad. Sci. USA*, 84:214-218
25 (1987), for example.

Nucleic acid molecules encoding the antibodies, monoclonal antibodies,

biologically active fragments, chimeric antibodies and CDR grafted antibodies described above also are provided by this invention. "Nucleic acid" is intended to include single and double stranded DNA, cDNA and RNA. These nucleic acid molecules can be operationally linked to promoter of RNA transcription.

5 The invention also encompasses nucleic acids molecules which differ from that of the nucleic acid molecules described above, but which produce the same phenotypic effect. The invention encompasses nucleic acid molecules characterized by changes in non-coding regions that do not alter the phenotype of the polypeptide produced therefrom when compared to the

10 nucleic acid molecule described hereinabove. This invention further encompasses nucleic acid molecules which hybridize to the nucleic acid molecule of the subject invention. As used herein, the term "nucleic acid" encompasses RNA as well as single and double stranded DNA and cDNA.

In one embodiment, these nucleic acid molecules are inserted into expression

15 vectors as noted above. The expression vectors can be inserted into suitable host cells. When the cells are induced to grow under conditions favoring transcription and translation of the inserted nucleic acid sequence, a recombinant protein or polypeptide is produced which can then be isolated and used for diagnosis or therapy as described below. Methods of

20 recombinantly producing polypeptides and proteins are generally known (see Sambrook, *et al.*, *supra* and Kreigler, M., *Gene Transfer and Expression: A Laboratory Manual*, W.H. Freeman and Company, New York, 1990, each incorporated herein by reference).

Pharmaceutical compositions also are provided by this invention. These

25 pharmaceutical compositions contain any of the above described polypeptides, fragments, antibodies, monoclonal antibodies, antibody fragments, chimeric antibodies or CDR grafted antibodies, each alone or in combination with each other, and a pharmaceutically acceptable carrier. As used herein, the term

"pharmaceutically acceptable carrier" encompasses any of the standard pharmaceutical carriers, such as a phosphate buffered saline solution, water, and emulsions, such as an oil/water or water/oil emulsion, and various types of wetting agents. These pharmaceutical compositions are useful for diagnostic or therapeutic purposes.

The monoclonal antibodies of the invention are suited for *in vitro* for use, for example, in immunoassays in which they can be utilized in liquid phase or bound to a solid phase carrier. In addition, the monoclonal antibodies in these immunoassays can be detectably labeled in various ways. Examples of types of immunoassays which can utilize monoclonal antibodies of the invention are competitive and non-competitive immunoassays in either a direct or indirect format. Examples of such immunoassays are the radioimmunoassay (RIA) and the sandwich (immunometric) assay. Detection of the antigens using the monoclonal antibodies of the invention can be done utilizing immunoassays which are run in either the forward, reverse, or simultaneous modes, including competition immunoassays and immunohistochemical assays on physiological samples. Those of skill in the art will know, or can readily discern, other immunoassay formats without undue experimentation.

The monoclonal antibodies of the invention can be bound to many different carriers and used to detect CD14. Examples of well-known carriers include glass, polystyrene, polypropylene, polyethylene, dextran, nylon, amyloses, natural and modified celluloses, polyacrylamides, agaroses, and magnetite. The nature of the carrier can be either soluble or insoluble for purposes of the invention. Those skilled in the art will know of other suitable carriers for binding monoclonal antibodies, or will be able to ascertain such, using routine experimentation.

There are many different labels and methods of labeling known to those of ordinary skill in the art. Examples of the types of labels which can be used in the present invention include enzymes, radioisotopes, fluorescent compounds, colloidal metals, chemiluminescent compounds, and bio-luminescent compounds. Those of ordinary skill in the art will know of other suitable labels for binding to the monoclonal antibodies of the invention, or will be able to ascertain such, using routine experimentation. Furthermore, the binding of these labels to the monoclonal antibodies of the invention can be done using standard techniques common to those of ordinary skill in the art.

Further provided herein is a method of blocking the binding of LPS/LBP complex to a CD14 receptor on the surface of a cell by contacting the cell with a monoclonal antibody capable of binding to the complex, e.g., a monoclonal antibody with the specificity and affinity of 28C5, or 23G4 or a biologically active fragment of the monoclonal antibody. Also disclosed is a method of inhibiting NF- κ B activation of a cell expressing CD14 receptor in the presence of a ligand (such as LPS or LPS/LBP) which is capable of inducing NF- κ B activation. This method provides contacting the cell with an antibody having the ability to specifically bind cell surface receptor CD14 and full length human soluble CD14 or a biologically active fragment of the antibody. Specific examples of such antibodies are the antibodies designated 28C5, 23G4 and 18E12.

Several therapeutic methods are provided herein which can be utilized in animals and humans. One therapeutic method is a means to treat or prevent disease associated with NF- κ B activation, such as sepsis, by administering to a subject with the disease an effective amount of an antibody having the ability to specifically bind cell surface receptor CD14 and soluble CD14 or a biologically active fragment of the antibody. The above method is especially advantageous when a monoclonal antibody having the binding specificity of

18E12 is used. Because 18E12 and the like antibodies can inhibit NF- κ B activation even after LPS has bound the CD14 receptor, such antibody can be used for the treatment of later stage sepsis. As used herein, later stage sepsis means the disease course after LPS has bound CD14 cell-associated receptor.

5 Significantly, 18E12 and like antibodies are capable of allowing the cell to which the antibody has bound to continue to transport LPS or LPS/LBP complex into the cell. This property provides the added benefit of allowing the removal of LPS or LPS/LBP complex from the *in vivo* system thereby inhibiting the possible pathological interaction of LPS or LPS/LBP complex at some other *in*

10 *vivo* site.

Alternatively, monoclonal antibodies 28C5 and 23G4 are preferred in the method of the invention where treatment is prophylactic or it is desirable to block LPS/LBP from binding to CD14, thereby inhibiting cytokine release and cell activation.

15 The invention provides a therapeutic method of ameliorating sepsis or one or more of the symptoms of sepsis comprising administering to a subject displaying symptoms of sepsis or at risk for developing sepsis, a therapeutically effective amount of a monoclonal antibody of the invention that binds to CD14 and inhibits cell activation. Such symptoms which may be

20 ameliorated include those associated with a transient increase in the blood level of TNF, such as fever, hypotension, neutropenia, leukopenia, thrombocytopenia, disseminated intravascular coagulation, adult respiratory distress syndrome, shock and multiple organ failure. Patients who require such treatment include those at risk for or those suffering from toxemia, such as endotoxemia resulting

25 from a gram-negative bacterial infection, venom poisoning, or hepatic failure, for example. In addition, patients having a gram-positive bacterial, viral or fungal infection may display symptoms of sepsis and may benefit from such a therapeutic method as described herein. Those patients who are more

particularly able to benefit from the method of the invention are those suffering from infection by *E. coli*, *Haemophilus influenza B*, *Neisseria meningitides*, staphylococci, or pneumococci. Patients at risk for sepsis include those suffering from burns, gunshot wounds, renal or hepatic failure.

5 The term "therapeutically effective amount" as used herein refers to the amount of monoclonal antibody which binds to CD14 and blocks signalling events such as cytokine release, used is of sufficient quantity to decrease the subject's response to LPS and decrease the symptoms of sepsis. The term "therapeutically effective" therefore includes that amount of antibody sufficient
10 to prevent, and preferably reduce by at least 50%, and more preferably sufficient to reduce by 90%, a clinically significant increase in the plasma level of TNF, for example. The dosage ranges for the administration of the monoclonal antibody of the invention, for example 18E12, 28C5 and 23G4, are those large enough to produce the desired effect. Generally, the dosage will
15 vary with the age, condition, sex, and extent of the infection with bacteria or other agent as described above, in the patient and can be determined by one skilled in the art. The dosage can be adjusted by the individual physician in the event of any contraindications. In any event, the effectiveness of treatment can be determined by monitoring the level of LPS and TNF in a patient. A
20 decrease in serum LPS and TNF levels should correlate with recovery of the patient.

In addition, patients at risk for or exhibiting the symptoms of sepsis can be treated by the method as described above, further comprising administering, substantially simultaneously with the therapeutic administration of a monoclonal
25 antibody of the invention, an inhibitor of TNF, an antibiotic, or both. For example, intervention in the role of TNF in sepsis, either directly or indirectly, such as by use of an anti-TNF antibody and/or a TNF antagonist, can prevent or ameliorate the symptoms of sepsis. Particularly preferred is the use of an

anti-TNF antibody as an active ingredient, such as a monoclonal antibody with TNF specificity as described by Tracey, *et al.* (*Nature*, 330:662, 1987).

5 A patient who exhibits the symptoms of sepsis may be treated with an antibiotic in addition to the treatment with a truncated LBP or antibody of the invention. Typical antibiotics include an aminoglycoside, such as gentamycin or a beta-lactam such as penicillin, or cephalosporin. Therefore, a preferred therapeutic method of the invention includes administering a therapeutically effective amount of an antibody of the invention, substantially simultaneously with administration of a bactericidal amount of an antibiotic.

10 The term "bactericidal amount" as used herein refers to an amount sufficient to achieve a bacteria-killing blood concentration in the patient receiving the treatment. The bactericidal amount of antibiotic generally recognized as safe for administration to a human is well known in the art, and as is known in the art, varies with the specific antibiotic and the type of bacterial infection being
15 treated.

Preferably, administration of a monoclonal antibody of the invention occurs within about 48 hours and preferably within about 2-8 hours, and most preferably, substantially concurrently with administration of the antibiotic.

20 For the purposes of this invention, a subject is an animal or a human patient and an effective amount is from about 0.25 mg/kg/body weight to about 50 mg/kg/body weight. In one embodiment, the effective amount is from about 0.5 mg/kg/body weight to about 10 mg/kg/body weight. When the subject is a human patient, the preferred amount is from about 0.5 mg/kg/body weight to about 8 mg/kg/body weight.

As is known to those of skill in the art, the above methods may be combined to enhance the therapeutic and prophylactic effects. Means of administering pharmaceutical compositions are well known to those of skill in the art and include, but are not limited to administration intravenously, orally, intraperitoneally, subcutaneously or by inhalation therapy.

The following examples are intended to illustrate but not limit the invention.

EXAMPLE 1

GENERATION OF SOLUBLE CD14 (sCD14) AND PRODUCTION OF MONOCLONAL ANTIBODIES

10 A. Cloning of sCD14

A copy of the human CD14 gene was obtained. The description of the cloning of this gene, from a human monocytic cell line (HL-60) (American Type Culture Collection, ATCC No. 240), is described in *Blood*, 73:284 (1989), incorporated herein by reference. The CD14 gene was excised from this expression vector and cloned into the mammalian expression vector pEE14 (Celltech). This vector has an inducible glutamine synthetase gene (GS) which was used to amplify the inserted DNA fragments containing the CD14 gene. A full-length DNA sequence of the gene was cloned into pEE14. Cells expressing soluble CD14 were identified as an ELISA assay by reactivity with commercially available anti-CD14 mAbs. One clone, identified as 523, was demonstrated to express both soluble CD14 and a membrane associated form which could be detected by FACS analysis. The soluble form of clone 523 was determined to be N-terminally processed at amino acid residue 20 of the predicted translated protein sequence. The sequence for this protein is set forth in Figure 1. Amino acid residues 1-19 of the translated CD14 sequence was predicted to be a signal sequence (Gene Works, Intelligenetics). It was determined by C-

terminal sequence analysis that the C-terminus was intact; no processing had occurred which was similar to that noted in the soluble CD14 isolated from human serum (Bazil, *et al.*, *Eur. J. Immunol.*, 16:1583, 1986, incorporated herein by reference). The soluble CD14 isolated from urine of nephritic patients
5 is lacking the eight most C-terminal amino acids (Bazil, *Mol. Immunol.*, 26:657, 1989). The clone 523 may have avoided the processing steps at the C-terminus as a consequence of its expression in CHO cells.

Purification of the soluble CD14 was accomplished by purifying the antigen on an affinity column composed of commercially available mAb 63D3 (obtainable
10 from the American Type Culture Collection (ATCC No. HB 44)).

B. Generation of sCD14 Monoclonal Antibodies

Monoclonal antibodies to human soluble CD14 were generated by somatic cell fusion between spleen cells from BALB/c mice immunized with purified human recombinants CD14 described above, and the mouse myeloma cell line
15 X63.Ag8.653. Monoclonal antibodies 28C5, 18E12, 26F3 and 23G4 are IgG1 mAb which were identified by screening against CD14 in an ELISA assay. Binding to native CD14 was confirmed by flow microfluoremetry on CD14⁺ cells and immunoprecipitation of biosynthetically labeled CD14. Monoclonal antibodies 28C5, 23G4 and 18E12 recognize cell-associated and soluble CD14.
20 Competition studies indicated that these mAb bound to three distinct CD14 epitopes (overlap between 28C5 and 23G4).

EXAMPLE 2
CHARACTERIZATION OF MONOCLONAL ANTIBODIES

A. **FACS Analysis of CD14⁺ Positive Cells**
with anti-CD14 Monoclonal Antibodies

5 THP-1 cells (American Type Culture Collection, ATCC No. TIB 202) were induced with dihydroxyvitamin D3 for 48 hours and cells were then washed with DMEM (Dulbecco's modified Eagles medium with 4.5 g/L glucose, 90% fetal bovine serum, 10% followed by PBS with 1% BSA and 0.02% azide). One million cells/tube were reacted with first antibody (anti-CD14s supernatants) at
10 1:2 dilution for 45 minutes at 4°C. Cells are washed with PBS/BSA/azide. The second antibody was added (goat anti-mouse IgG, FITC-labeled (Cappel)) at a 1:250 dilution for 30 minutes at 4°C. Cells are washed 2X with same buffer. Pellets resuspended in 1ml of buffer. Fluorescence intensity was measured by
15 a Cytofluorograph (Ortho Instruments). The results are shown in Figures 6 through 10.

B. **Determination of Binding Affinity**
of Anti-sCD14 Monoclonal Antibodies for Soluble CD14

All anti-CD14 mAbs, and some commercially available anti-CD14s, were evaluated for antibody affinity to soluble CD14 antigen (Figure 11). Antibody
20 3C10 had a titer of 1:2⁸ and 28CS a titer of 1:2¹¹. This shows a eight-fold difference in titer of 28C5 as compared to 3C10. Relative affinities of the antibodies were determined at equivalent concentrations of purified antibody protein which were probed with a labeled goat anti-mouse conjugated antibody. Those antibodies with the highest affinity for sCD14, presented in
25 this manner, are 4F2, 5G3, 26F3, 28C5, 23G4 and 63D3. Anti-CD14 mAbs 3C10 and 18E12 exhibited the lowest affinity for sCD14. Anti-CD14 28C5 had

a much higher affinity as compared to 3C10. Soluble CD14 was coated onto microtiter plates and the anti-CD14 mAbs added in two-fold serial dilutions starting with 2 μ g/ml. A goat anti-mouse HRP conjugated antibody is added and the plates incubated and washed. Substrate is then added and the 1:2ⁿ titer is recorded. This is represented by the highest dilution of antibody giving at least 3X the OD_{490nm} value of a negative control.

Competition studies between labeled 3C10 and a panel of anti-CD14 mAbs revealed that only antibodies 28C5 and 10A1 (an IgA mAb) were capable of competing with 3C10 for sCD14 on the coated microtiter plate (Figure 12). Further competition assays were performed and confirmed that 18E12 did not compete with 28C5, 3C10, or 26F3.

Table 1 shows the results of a similar competition assay between 28C5, 23G4 and 18E12. The antibodies were coated on the solid phase in an ELISA assay with biotin labeled anti-CD14 mAbs (A,C).

TABLE 1
COMPETITION STUDIES BETWEEN DIFFERENT ANTI-CD14
MABS FOR CD14 RECOGNITION AND THE ABILITY OF THESE
MABS TO BLOCK LPS/LBP BINDING TO CD14

	A	B	C	D
Antibody	+18E12	+28C5	+23G4	LPS Binding
	% Inhibition	% inhibition	% inhibition	% inhibition
18E12	95.5	59.8	30.4	3.6
28C5	46.1	90.6	69.7	83.3
23G4	77.3	95.5	95.5	85.4

The ability of these anti-CD14 mAbs to block LPS/LBP binding to sCD14 was assessed using as similar ELISA format.

When the mAbs were evaluated for ability to inhibit LPS/LBP binding to sCD14, 28C5 was the most effective (Figure 13). Figure 13 represents the intensity of binding of LPB/biotinylated LPS complex to soluble CD14 immobilized on a solid phase, in the presence of 4 μ g/ml of different anti-CD14 monoclonal antibodies. Anti-CD14 mAb 28C5 and 3C10 block this binding event. Anti-CD14 mAb is more efficient in its blocking as noted by the decreased OD value. Anti-CD14 mAb 18E12 does not exhibit any blocking effect.

Evaluation of the anti-CD14 mAbs for the ability to block cytokine release in HL-60 cells in response to LPS stimulation, showed that 28C5 blocked TNF- α expression (Figure 14). Inhibition of cytokine release was also observed when 28C5, 23G4, and 18E12 were added to whole blood *ex vivo*

prior to addition of LPS. Surprisingly, 18E12 inhibited cytokine release even though it was previously demonstrated not to block LPS/LBP binding to CD14 (Figure 13 and 27). The effect of 23G4, 28C5 and 18E12 on LPS-induced TNF in baboon whole blood was also examined, *ex vivo*. The results in Figure 28 show that 23G4 was most effective at inhibiting TNF secretion in LPS-induced baboon blood. These results show that 18E12 was specific for a domain on CD14 which did not prevent LPS/LBP binding, but is important to the signaling events involved in LPS stimulation of cells.

Although 28C5 and 23G4 share specificity in blocking binding of LPS:LBP to CD14, they do not share recognition of baboon CD14 (only 23G4 recognizes baboon), nor can 28C5 block TNF α release from baboon whole blood in response to LPS (Figure 27).

The ability of the anti-CD14 mAbs, 28C5, 18E12 and 23G4 to block LPS/LBP binding to sCD14 was also assessed using a similar ELISA format as described for Table 1. The results also show that 23G4 and 28C5 compete for sCD14 binding (see Table 1).

C. Activation of Cytokine Release

HL-60 cells (obtained from the American Type Culture Collection, ATCC No. CCL-240) were plated at a concentration of 1.5×10^5 cells per ml. The cells were induced toward the monocyte lineage for 3 to 4 days in RPMI 1640 containing 10% bovine serum, 10^{-7} M DHvD3 (Biomol Research Laboratories) and 50 μ M indomethacin (Calbiochem). These differentiated cells were resuspended at 1×10^6 cells/ml growth medium containing 50 μ M indomethacin with or without 10% human type AB serum (Irvine Scientific) and then were added to flat bottom cluster dishes. Cells were activated by addition of different concentrations of LPS (*E. coli* serotype 01217:B8; Sigma) followed by a 4 to 5 hour incubation at 37°C. Cells in the culture

plates were pelleted by low speed centrifugation (170 X g for 10 minutes at room temperature) and the growth medium was removed for ELISA (ELISA kit for human TNF- α detection; Genzyme) of soluble cytokine levels.

D. Inhibition of LPS binding to Cellular CD14 by Anti-CD14 Monoclonal Antibodies

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To characterize the mechanism of interaction between cellular CD14 and LPS, stably transfected 70Z/3 cells containing the human CD14 expression vector described in Lee, *et al.*, *J. Exp. Med.*, 175:1697-1705, 1992, were prepared to form 70Z/3-hCD14 cells. Stably transfected cells expressing cell surface CD14 were confirmed using FACS analysis on cells stained with FITC conjugated anti-human CD14 Mab MY4 described earlier. Other transfected cell lines were also prepared expressing CD14 fusion proteins in which the membrane anchor from CD14 was removed and replaced with the membrane anchor from decay accelerating factor (DAF), designated 70Z/3-hCD14DAF, with the membrane anchor from human tissue factor, designated 70Z/3-hCD14TF, and with the membrane anchor from the murine class molecule, H2K², designated 70Z/3-hCD14CI.

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Direct binding of LPS to cellular CD14 was characterized using FITC-labelled LPS (FITC-LPS). 70Z/3-CD14 cells were suspended in culture medium containing 10% FCS with or without 10 ug/ml Mab, and pre-incubated for 30 minutes at 37°C. Thereafter, FITC-Re595-LPS was added at 1 ng/ml and maintained for 15 minutes at 37°C. Immediately thereafter, an equal volume of ice-cold RPMI 1640 medium was added, and the admixture was maintained at 4°C until FACS analysis. Cell-associated fluorescence was measured as described by Lee *et al.*, supra, and measured fluorescence adjusted by subtracting fluorescence measured using non-transfected 70Z/3 cells.

The bar-graph results are shown in Figure 15, and indicate that for all transfected cell types that contain a cell-surface CD14 protein (wt or fusion protein), the anti-CD14 monoclonal antibody Mab MY4 (shaded bar) blocked LPS binding to cells, whereas Mab 18E12 (open bar) did not block LPS binding to cells. Binding of FITC-LPS in the presence of Mab 18E12 was similar to results obtained using no antibody (black bar). Mab MY4 is an antibody known to immunoreact with CD14, and by the data presented herein is shown to inhibit LPS binding to CD14 and to inhibit LPS-dependent, CD14-mediated cell activation. The differences in levels of FITC-LPS binding reflect the differences in levels of CD14 expression in the different transfected cell lines. Whereas the hCD14 transfected cells contain approximately 10,000 receptors per cell, the hCD14DAF transfected cells contain approximately 50,000 receptors per cell, and the hCD14Tf and hCD14Cl transfected cells are estimated to each contain about 15,000-20,000 receptors per cell. The results are expressed as a mean +/- standard deviation of three independent determinations.

E. Inhibition of LPS-Dependent, CD14-Mediated Activation of Cells using Anti-CD14 Monoclonal Antibodies

Anti-CD14 monoclonal antibodies were characterized for their ability to inhibit CD14-mediated activation of cells by LPS. To that end, a CD14 transfected cell system was developed and demonstrated to be responsive to LPS-induced activation. Several transfected 70Z/3 cell lines were prepared as described in Example 3, and contain several membrane associated forms of CD14 as described earlier.

The transfected cells were cultured as described by Lee et al., *supra*, suspended in RPMI 1640 media containing 10% fetal calf serum (FCS; heat inactivated, 56°C for 30 min.) and 10 ug/ml antibody (MY4 or 18E12) as

indicated by a "+" in Figure 16, and maintained for 30 min at 37°C. Thereafter, 100uM taxol or LPS (1 ng/ml Re595 LPS) was added as indicated by a "+" in Figure 16 and the cells were maintained for 15 min at 37°C. Thereafter, the cells were harvested and nuclear extracts was
5 prepared to determine activation of NF- κ B as described by Molitor *et al.*, *Proc. Natl. Acad. Sci. USA*, 87:10028-10032 (1990). P³²-labelled NF- κ B-specific oligonucleotides (5'-CAGAGGGGACTTTCCGAGA-3') in double-stranded form were used in a gel retardation assay to detect the presence of NF- κ B on 4% non-denaturing polyacrylamide gels.

10 The results of the study are shown in Figure 16, and indicate that LPS and taxol both induce NF- κ B activation. As expected, LPS induced minimal NF- κ B activation in control transfected cells lacking CD14 (70Z/3-RSV), and induced marked activation in cells expressing CD14. This shows that LPS-induced NF- κ B activation is mediated by and requires CD14 on the cell
15 surface. Furthermore, the results show that both MY4 and 18E12 inhibit LPS-induced NF- κ B expression, but not taxol-induced NF- κ B expression, indicating that the inhibitory effect of the antibodies is specific and dependent upon CD14.

20 These results with Mabs MY4 and 18E12 indicate that LPS binding to CD14 is not sufficient to induce cell activation and that additional interactions following LPS-CD14 binding are critical for cell activation. The results also indicate that inhibition of CD14-mediated activation of cells may occur at different levels, first by blocking the inducer (LPS) from binding to CD14, and second by blocking a subsequent step after inducer binds to CD14.
25 The data also establish that the use of inhibitors of the second step will block CD14-mediated cell activation where the inducer is a molecule other than LPS.

F. Inhibition of LPS Uptake by CD14⁺ Cells

In the progression of sepsis, LPS binds cell surface CD14, and is known to be taken up by those CD14⁺ cells (Kitchens, *et al.*, *J. Exp. Med.*, 176:485-494, 1992; Pugin, *et al.*, *PNAS*, 90:2744-2748, 1993). The difference
5 between antibodies that block LPS binding to CD14, such as Mab 28C5, and antibodies that do not block binding to CD14, such as Mab 18E12, is significant in view of LPS uptake by CD14⁺ cells. Therefore, the ability of anti-CD14 antibodies to inhibit LPS uptake was characterized.

To that end, FITC-LPS uptake on transfected 70Z/3-hCD14 cells was
10 measured in the presence of various anti-CD14 antibodies. Fluorescence arising from the FITC-LPS was detected inside CD14⁺ transfected cells when no antibody was utilized after the cells were maintained at 37°C following exposure to FITC-LPS, confirming that LPS is taken up by cells under normal conditions. In the presence of Mab 28C5, uptake of LPS
15 could be inhibited completely, whereas in the presence of Mab 18E12, uptake could only be reduced to about 65% of the amount of uptake observed under normal conditions. These results indicate that Mab 18E12 is particularly useful for inhibiting CD14⁺ cell activation where it is desirable to allow LPS to enter the cells, because Mab 18E12 does not substantially
20 prevent LPS uptake. Subsequent studies show that in the presence of Mab 23G4, uptake of LPS could be inhibited similar to that seen with Mab 28C5.

CD14 Antigen Assay (ELISA)

- Coating: 150 μ l/well of anti-CD14 mAb 28C5 diluted at 1 μ g/ml in bicarbonate buffer. Incubate overnight at 4°C.
- 5 Blocking: Wash the plate 4X then add 150 μ l/well of blocking buffer. Incubate 1 hr. at 37°C.
- Samples: Wash the plate 1X then add 125 μ l/well of samples diluted in dilution buffer. Incubate 1 hr. at 37°C.
- 10 Conjugate: Wash the plate 5X then add 0.100 ml/well of biotinylated anti-CD14 mAb 18E12 diluted at 1 μ g/ml in dilution buffer. Incubate 1 hr. at 37°C.
- 15 Av-HRPO: Wash the plate 5X then add 0.100 ml/well of preformed streptavidin/biotin/peroxidase complex. (Streptavidin/biotinylated/HRPO preparation (Zymed SABC kit): Mix 2 μ l/ml of Streptavidin with 2 μ l/ml of biotinylated-HRPO in washing buffer and incubate 30 minutes at 37°C. Before adding to the wells, dilute at 1:2 with dilution buffer.) Incubate 30 minutes at 37°C.
- 20 Substrate: Wash the plate 5X then add 0.100 ml/well of Sigma OPD, leave the plate 30 minutes in the dark and stop the color development with 0.050 ml of 4N H₂SO₄. Read plate at 490 nm.
- CD14 Standard: 2-fold serial dilutions of clone 523 at 100 ng/ml.
- Serum Dilutions: Starting dilution 1:25-1:50.

Miscellaneous ELISA Reagents

Blocking Buffer: PBS + 10% w/v of nonfat dry milk (Carnation).

Washing Buffer: PBS + 0.05% v/v of Tween 20.

Dilution Buffer: Mix vol/vol blocking buffer and wash buffer, use to dilute samples, labelled antibody and the preformed complex.

EXAMPLE 3

MONOCLONAL ANTIBODY CLONING

Messenger RNA was extracted from monoclonal antibody producing cell lines using the method of Chomczynski and Sacchi, *Anal. Bio.*, 162:156-159 (1987), incorporated herein by reference. Reverse transcription was performed using murine specific 3' antibody primers (IgG1 or k) and the resulting cDNAs subjected to PCR (Supplier) according to the manufacturer's instructions, using a panel of murine specific 5' antibody primers described in Huse, *et al.*, *Science*, 246:1275-1281 (1989), incorporated herein by reference. Heavy and light chain DNA fragments were gel purified and digested with appropriate enzymes. The 672 base pair heavy chain fragment was cloned into the Spe1/Xho1 site of pBluescript II KS⁺ and sequenced using the automated ABI Model 373A DNA sequencer, according to the manufacturer's instructions. The 642 base pair light chain fragment was cloned into the Sst1/Xba of pBluescript II KS⁺ and sequenced in a similar manner.

SEQ ID NO:1 and 2 are the nucleotide and deduced amino acid sequence of the 28C5 heavy chain and SEQ ID NO:3 and 4 are the nucleotide and deduced amino acid sequence of the 28C5 light chain. SEQ ID NO:5 and 6 are the the nucleotide and deduced amino acid sequence of the 18E12 heavy chain and SEQ ID NO:7 and 8 are the nucleotide and deduced amino acid sequence of the 18E12 light chain (Figures 2-5). Figure 30 shows a comparison of the heavy chains of 3C10, 28C5 and 18E12. SEQ ID NO:25 and 26 are the nucleotide and deduced amino acid sequence of the 23G4 light chain (See Figure 29). Figure 29 shows the amino acid sequence of the light chains of monoclonal antibodies 3C10, 28C5, 23G4 and 18E12. Figure 30 shows the amino acid sequence of the heavy chains of

monoclonal antibodies 3C10, 28C5, and 18E12.

Note that although both 28C5 and 23G4 share the same specificity in that they block LPS:LBP binding to CD14, compete with each other for sCD14 binding and prevent TNF α release in human whole blood at similar concentrations (see Figure 27), their light chains do not share the same nucleotide and amino acid sequence. (See Table 2)

Recombinant Expression of Nucleic Acids

The recombinant expression of nucleic acids of this invention are performed according to the following general strategy. PolyA⁺ mRNA is isolated from the antibody-expressing hybridoma cells. cDNA synthesis and PCR amplification of the mRNA are performed by methods described above. From the cDNA sequence data obtained, the amino acid sequences of the polypeptides encoded by the DNA sequences are deduced by a computer software program, for example, MAPSEQ, commercially available from DNASTar (Madison, Wisconsin).

The expression products, assembled as an antibody fragment, are screened for binding affinity by methods known in the art such as ELISAs (Enzyme-Linked Immuno-Sorbent Assay) utilizing the hapten or antigen, or affinity columns (as described, for example, in Skerra and Pluckthun, *Science*, 240:1038-1041, 1988, incorporated herein by reference).

Several types of vectors are available and can be used to practice this invention, e.g., plasmid, DNA and RNA viral vectors, baculoviral vectors, and vectors for use in yeast. When the vector is a plasmid, it generally contains a variety of components including promoters, signal sequences, phenotypic selection genes, origin of replication sites, and other necessary components as are known to those of skill in the art.

Promoters most commonly used in prokaryotic vectors include the *lac Z* promoter system, the alkaline phosphatase *pho A* promoter, the bacteriophage λ PL promoter (a temperature sensitive promoter), the *tac* promoter (a hybrid *trp-lac* promoter that is regulated by the *lac* repressor),
5 the tryptophan promoter, and the bacteriophage T7 promoter.

Promoters used to practice this invention are the *lac Z* promoter and the *pho A* promoter. The *lac Z* promoter is regulated by the *lac* repressor protein *lac i*, and thus transcription of the polypeptide can be controlled by manipulation of the level of the *lac* repressor protein. By way of illustration,
10 a phagemid containing the *lac Z* promoter is grown in a cell strain that contains a copy of the *lac i* repressor gene, a repressor for the *lac Z* promoter. Exemplary cell strains containing the *lac i* gene include JM 101 and XL1-blue. In the alternative, the host cell can be cotransfected with a plasmid containing both the repressor *lac i* and the *lac Z* promoter.
15 Occasionally both of the above techniques are used simultaneously, that is, phagemid particles containing the *lac Z* promoter are grown in cell strains containing the *lac i* gene and the cell strains are cotransfected with a plasmid containing both the *lac Z* and *lac i* genes. Normally when one wishes to express a gene, to the transfected host above, one would add an
20 inducer such as isopropylthiogalactoside (IPTG), but this step can be omitted.

Another useful component of vectors used to practice this invention is a signal sequence. This sequence is typically located immediately 5' to the gene encoding the polypeptide, and thus will be transcribed at the amino
25 terminus of the fusion protein. However, in certain cases, the signal sequence has been demonstrated to be located at positions other than 5' to the gene encoding the protein to be secreted. This sequence targets the protein to which it is attached across the inner membrane of the bacterial

- cell. The DNA encoding the signal sequence can be obtained as a restriction endonuclease fragment from any gene encoding a protein that has a signal sequence. Suitable prokaryotic signal sequences can be obtained from genes encoding, for example, LamB or OmpF (Wong, *et al.*, *Gene*, 68:193, 1983, incorporated herein by reference), MalE, PhoA, OmpA and other genes. A preferred prokaryotic signal sequence for practicing this invention is the *E. coli* heat-stable enterotoxin II (STII) signal sequence as described by Chang, *et al.*, *Gene*, 55:189, 1987, incorporated herein by reference.
- 10 Another useful component of the vectors used to practice this invention is a phenotypic selection gene. Typical phenotypic selection genes are those encoding proteins that confer antibiotic resistance upon the host cell. By way of illustration, the ampicillin resistance gene (*amp*), and the tetracycline resistance gene (*tet*) are readily employed for this purpose.
- 15 Construction of suitable vectors comprising the aforementioned components as well as the gene encoding the desired polypeptide are prepared using standard recombinant DNA procedures. References for recombinant methodology have been provided *infra*. Isolated DNA fragments to be combined to form the vector are cleaved, tailored, and ligated together in a
- 20 specific order and orientation to generate the desired vector.
- The DNA is cleaved using the appropriate restriction enzyme or enzymes in a suitable buffer. In general, about 0.2-1 μ g of plasmid or DNA fragments is used with about 1-2 units of the appropriate restriction enzyme in about 20 μ l of buffer solution. Appropriate buffers, DNA concentrations, and
- 25 incubation times and temperatures are specified by the manufacturers of the restriction enzymes. Generally, incubation times of about one or two hours at 37°C are adequate, although several enzymes require higher

temperatures. After incubation, the enzymes and other contaminants are removed by extraction of the digestion solution with a mixture of phenol and chloroform and the DNA is recovered from the aqueous fraction by precipitation with ethanol.

- 5 To ligate the DNA fragments together to form a functional vector, the ends of the DNA fragments must be compatible with each other. In some cases, the ends will be directly compatible after endonuclease digestion. However, it may be necessary to first convert the sticky ends commonly produced by endonuclease digestion to blunt ends to make them compatible for ligation.
- 10 To blunt the ends, the DNA is treated in a suitable buffer for at least 15 minutes at 15°C with 10 units of the Klenow fragment of DNA polymerase I (Klenow) in the presence of the four deoxynucleotide triphosphates. The DNA is then purified by phenol-chloroform extraction and ethanol precipitation.
- 15 The cleaved DNA fragments are size-separated and selected using DNA gel electrophoresis. The DNA is electrophoresed through either an agarose or a polyacrylamide matrix. The selection of the matrix will depend on the size of the DNA fragments to be separated. After electrophoresis, the DNA is extracted from the matrix by electroelution, or, if low-melting agarose is used
- 20 as the matrix, by melting the agarose and extracting the DNA from it.

- The DNA fragments that are to be ligated together (previously digested with the appropriate restriction enzymes such that the ends of each fragment to be ligated are compatible) are put in solution in about equimolar amounts. The solution will also contain ATP, ligase buffer and a ligase such as T4
- 25 DNA ligase at about 10 units per 0.5 µg of DNA. If the DNA fragment is to be ligated into a vector, the vector is at first linearized by cutting with the appropriate restriction endonuclease(s). The linearized vector can then be

treated with alkaline phosphatase or calf intestinal phosphatase. The phosphatasing prevents self-ligation of the vector during the ligation step.

After ligation, the vector with the foreign gene now inserted is transformed into a suitable host cell. Suitable prokaryotic host cells include *E. coli* strain
5 JM101, *E. coli* K12 strain 294 (ATCC number 31,446), *E. coli* strain W3110 (ATCC number 27,325), *E. coli* X1776 (ATCC number 31,537), *E. coli* XL-1Blue (Stratagene), and *E. coli* B; however, many other strains of *E. coli*, such as HB101, NM522, NM538, NM539 and many other species and genera of prokaryotes can be used as well. In addition to the *E. coli* strains
10 listed above, bacilli such as *Bacillus subtilis*, other enterobacteriaceae such as *Salmonella typhimurium* or *Serratia marcesans* and various *Pseudomonas* species can all be used as hosts.

Transformation of prokaryotic cells is readily accomplished using calcium chloride or other methods well known to those skilled in the art.
15 Electroporation (Neumann, *et al.*, *EMBO J.*, 1:841 1982, incorporated herein by reference) also can be used to transform these cells. The transformed cells are selected by growth on an antibiotic, commonly tetracycline (tet) or ampicillin (amp), to which they are rendered resistant due to the presence of tet and/or amp resistance genes on the vector.

20 After selection of the transformed cells, these cells are grown in culture and the plasmid DNA (or other vector with the foreign gene inserted) is then isolated. Plasmid DNA can be isolated using methods known in the art. This purified plasmid DNA is then analyzed by restriction mapping and/or DNA sequencing.

25 Following procedures outlined above, mammalian cell lines such as myeloma (P3-653), hybridoma (SP2/0), Chinese Hamster Ovary (CHO),

Green monkey kidney (COS1) and murine fibroblasts (L492) are suitable host cells for polypeptide expression. These "mammalian" vectors can include a promoter, an enhancer, a polyadenylation signal, signal sequences and genes encoding selectable markers such as geneticin (neomycin
5 resistance), mycophenolic acid (xanthine guanine phosphoribosyl transferase) or histidinol (histidinol dehydrogenase).

Suitable promoters for use in mammalian host cells include, but are not limited to, Ig Kappa, Ig Gamma, cytomegalovirus (CMV) immediate early, Rous Sarcoma Virus (RSV), simian virus 40 (SV40) early, mouse mammary
10 tumor (MMTV) virus and metallothionein. Suitable enhancers include, but are not limited to Ig Kappa, Ig Heavy, CMV early and SV40. Suitable polyadenylation sequences include Ig Kappa, Ig Gamma or SV40 large T antigen. Suitable signal sequences include Ig Kappa, Ig Heavy and human growth hormone (HGH).

15 When the vector is baculovirus, suitable promoters and enhancer sequences include, but are not limited to AcMNPV polyhedrin, AcMNPV ETL and AcMNPV p10 sequences. One particularly suitable polyadenylation signal is the polyhedrin AcMNPV. Ig Kappa, Ig Heavy and AcMNPV are examples of suitable signal sequences. These vectors are useful in the following insect
20 cell lines, among others: SF9, SF21 and High 5.

Alternatively, the polypeptides can be expressed in yeast strains such as PS23-6A, W301-18A, LL20, D234-3, INVSC1, INVSC2, YJJ337. Promoter and enhancer sequences such as gal 1 and pEFT-1 are useful. Vra-4 also provides a suitable enhancer sequence. Sequences useful as functional
25 "origins of replication" include ars1 and 2 μ circular plasmid.

TABLE 2
AMINO ACID SEQUENCE ANALYSIS OF ANTI-CD14 mAbs CDRs

	<u>MAb</u>	<u>CDR1</u>	<u>CDR2</u>	<u>CDR3</u>
5	3C10-Heavy	SYAMS (SEQ ID NO: 10)	SISSGGTTYYPDNVKG (SEQ ID NO: 11)	GYDYHY (SEQ ID NO: 12)
	28C5-Heavy	SDSAWN (SEQ ID NO: 13)	YISYSGSTSYNPSLKS (SEQ ID NO: 14)	GLRFAY (SEQ ID NO: 15)
10	% homology	17%	38%	14%
<hr/>				
15	3C10-Light	RASESVDSFGNSFMH (SEQ ID NO: 16)	RAANLES (SEQ ID NO: 17)	QQSYEDPWT (SEQ ID NO: 18)
	28C5-Light	RASESVDSYVNSFLH (SEQ ID NO: 19)	RASNLQS (SEQ ID NO: 20)	QQSNEDPYT (SEQ ID NO: 21)
20	23G4-Light	RASESVDSYGKSFMH (SEQ ID NO: 22)	VASKLES (SEQ ID NO: 23)	QQNNEDPYT (SEQ ID NO: 24)
	% homology	80%	71%	67%

EXAMPLE 4**IN VIVO TREATMENT WITH CD14 ANTIBODIES**

The pretreatment of rabbits with IFN- γ for three days followed by an injection of LPS produces a sepsis state in rabbits (G.J. Jurkovich, *et al.*, *J. Surg. Res.*, 51:197-203, 1991). A similar protocol was followed in the *in vivo* experiments described herein. 5 μ g/kg of IFN- γ (specific activity: 2.5×10^8 units/mg; 5 μ g = 1.25×10^6 units/kg) was injected daily for three consecutive days, then following establishment of baseline cardiac output and systemic pressure, an 8 hour infusion of LPS (3 mg/kg total dose or 375 μ g/kg/hour x 8 hours) was started. Subcutaneous injections of IFN- γ were given 3 days and on the 3rd day baseline data was collected for at least one hour prior to starting the infusion of LPS. These animals were maintained on ketamine throughout the experiment. The rabbit experiments suggested that these animals will become somnolent following the LPS infusion. The same established protocol was also utilized in two groups of monkeys. Animals were randomly assigned to either isotype matched control monoclonal antibodies (MAbs) or a CD14 blocking MAb with the person responsible for their care unaware of the treatment protocol. A dose of 5mg/kg/mAb (isotype or CD14-specific) was given by bolus injection 30' prior to the start of the LPS infusion.

Animals were anesthetized with ketamine then arterial and venous catheters placed in the femoral artery and vein, respectively. The arterial catheter has a thermistor at its tip for determination of thermal dilution cardiac output. A second lumen on this catheter was used for arterial pressure measurement. The venous catheter was used to infuse drugs, maintenance fluids and for cold injection in the cardiac output measurements. Lactated Ringers (3 ml) was used for each cardiac output determination.

- Blood pressure and cardiac output was recorded every 15 minutes throughout the baseline period and then every half hour for the remainder of the experiment. Blood was drawn (3 ml) every hour for determination of arterial PO_2/PCO_2 , pI and protein. This same blood sample was used for determination of systemic white blood cell counts and differential counts. These animals were resuscitated with lactated Ringers' solution following the infusion of LPS. All animals were given an infusion of 4mls/kg as a maintenance infusion and this was increased as necessary to maintain cardiac output to within 10% of baseline.
- A total of six animals each were pretreated with either the IgG1 isotype control or 28C5, and five animals were pretreated with 18E12. All test animals were challenged with LPS 30 minutes after the infusion of antibody. Seventy-two hours prior to LPS infusion, monkeys were given 3 subcutaneous injections of human recombinant interferon gamma (125,000 U/Kg) at 24 hour intervals. To measure MAP levels in anesthetized animals, arterial and venous catheters were placed in the femoral artery and vein, respectively as described above.

Mean Arterial Pressure (MAP) of Monkeys Challenged with LPS

- The mean arterial pressure (MAP) results reveal that pretreatment with 28C5 prevented a significant drop in blood pressure, particularly at the 2 hour time point, common in the control group (FIGURE 17). However, animals pretreated with 18E12 exhibited this drop in blood pressure at 2 hours, yet were able to recover to percents noted in the 28C5 pretreated animals. The function of 18E12 differs from that of 28C5 in that only signaling events are prevented, not inhibition of LPS binding; a key feature of 28C5 as well as 23G4. This difference in function may reflect the difference noted in the MAP response. Protection by 18E12 may involve late LPS-induced effects. This MAP profile by 18E12 suggests that even in the event of physiological

responsiveness to LPS (presence of hypotension) this anti-CD14 mAb is capable of preventing the deleterious effects noted in the isotype control-treated animals.

Effect of IFN On CD14 Concentration

- 5 The pretreatment of animals with interferon-gamma for three consecutive days, 24 hours apart, had relatively little effect on the CD14 concentration (FIGURE 18). However, the circulating levels of lipopolysaccharide binding protein (LBP) increased significantly (LBP was measured by ELISA in which
10 two non-competing monoclonal antibodies were used to capture and probe test samples); to levels noted during gram-negative sepsis. It is proposed that the interferon-gamma induces an acute phase response and sensitizes the animals to doses of LPS which otherwise would not induce any physiological and/or biochemical changes in these animals.

Lavage/Plasma Ratio of BSA

- 15 The Lavage/Plasma ratio of BSA is an indicator of lung damage and accesses the amount of BSA (injected one hour prior to the termination of the experiment) that permeates the lung tissue. The lung is one of the primary organs affected during endotoxemia. BSA levels were determined by an immunoassay utilizing a BSA specific monoclonal antibody. BSA
20 monoclonal antibodies are widely available. In this instance, the animals pretreated with 28C5 were protected significantly from the lung damage that was evident in the control-treated group (FIGURE 19). While 18E12-treated animals were not fully protected from the LPS effects, as a group they did better than the control animals.

Antibody Half-Life

25 A question arises as to the antibody half-life, or fate of the antibody due to its binding to both the soluble circulating form of CD14 and the membrane-

associated CD14 present on monocytes and neutrophils. When compared to an isotype control antibody which does not recognize human antigens, the kinetics of clearance are similar for all three groups (FIGURE 20).

CD14 Levels in Monkeys Treated With Antibody

5 CD14 was measured by ELISA in which two non-competing monoclonal antibodies are used to capture and probe test samples. The CD14 levels in monkeys treated with antibody only (no LPS challenge) were significantly higher in the 28C5 versus the control-treated animals (FIGURE 21). The reason for this rise is unknown although *in vitro* studies demonstrated that
10 exposure of CD14-bearing cells to anti-CD14 mAb 28C5 resulted in higher sCD14 levels; perhaps the antibody enhances the shedding mechanism of this GPI-linked protein. The 18E12-treated animals showed an increase at 36 hours which began to drop at 60 hours. When antibody-treated animals are challenged with LPS, there is no additional rise in CD14 levels in the
15 28C5 group, over what is noted in the antibody-only animal, suggesting that this is clearly an antibody-induced effect.

LBP Levels

The LBP levels rise after interferon-gamma treatment, as was mentioned above. In the anti-CD14-treated, LPS challenged animals there is a slight
20 lowering of detectable LBP levels versus control (FIGURE 22). The reason for this is not known although it may represent clearance of the complexes if the targets are unavailable for binding or transfer of the LPS.

ALT/GPT Levels

Enzyme transaminases ALT/GPT (considered the same enzyme) are
25 indicators of liver function and as such were measured to determine if there was evidence of necrosis. In patients with septic shock, the onset of hepatic failure is an early event in the MSOF syndrome. Maximum levels in

humans, depending upon the extent of damage, can reach 4000 U/ml. ALT/GPT levels were measured by following the manufacturer's recommendations in a test kit from Sigma Diagnostics.

5 While none of the levels recorded here are in the range noted in an extreme human situation, there is a trend in the control group for elevated enzyme levels during the 24 hour time course of the experiment. Normal levels for ALT/GPT in monkeys treated with antibody only, ranged from a mean of 18U/ml (T=0) to 52.2 U/ml (T=24), with the elevation at T=24 attributed to the anesthesia used throughout the study (ketamine). The two anti-CD14
10 treated groups followed a similar course; a mean of 16.5 U/ml at T=0 to 64.6 U/ml at T=24. The mean of the control group was 22.8 U/ml at T=0 to 98 U/ml at T=24. A twenty-four hour time course may not allow one to determine if the elevation in the control group of animals would continue (FIGURE 23).

15 Soluble E-Selectin Levels

Soluble E-selectin levels were measured by Parameter ELISA kit (British Biotechnology Products, Ltd.) to determine if blocking the CD14 receptor would somehow prohibit release of soluble E-selectin from endothelial cells. E-selectin expression on the surface of endothelial cells is an indicator of
20 activation of these cells and occurs as a consequence of TNF, IL-1 or LPS stimulation. Soluble E-selectin levels were elevated to similar levels, at 24 hours, in all groups (FIGURE 24).

IL-1, IL-6, IL-8 and TNF Levels

The cytokine response to LPS challenge was evaluated using Quantikine™
25 kits (R&D Systems), performed according to the manufacturer's specifications, in all groups of animals. The assay is an immunoassay with a solid phase ELISA format. The TNF α assay was a Biokine® enzyme

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immunoassay kit (T Cell Diagnostics) and was performed according to the manufacturer's specifications. It is known that $\text{TNF}\alpha$ and $\text{IL-1}\beta$ are key mediators of the inflammatory response induced as a consequence of LPS stimulation. In the anti-CD14 treated groups, the $\text{TNF}\alpha$ and $\text{IL-1}\beta$ responses were reduced versus the control treated group with 18E12 exhibiting the lowest level of expression of these inflammatory cytokines. Also, the peak TNF response was delayed by an hour in both anti-CD14 groups, the significance of this finding is not known at this time (FIGURE 25).

It is known that IL-1, IL-6 and IL-8 peak later than TNF in human septicemia, which is in line with the observations that release of IL-1, IL-6 and IL-8 is largely dependent on TNF generation. One cytokine which has been correlated with mortality in humans diagnosed with gram-negative sepsis is elevated IL-6 levels. IL-6 coordinates various aspects of the host defense against tissue injury. In the present model, 28C5 exhibited the lowest level of IL-6 in response to LPS. Animals pretreated with 18E12 also had lower levels than the control group, yet not as low as 28C5. The IL-8 response, while not significantly lower in the anti-CD14 groups, was reduced slightly (FIGURE 26). IL-8 levels in baboon models were shown to be correlated with TNF levels; reduced TNF resulted in reduced IL-8 levels. IL-8 has chemoattractant and granulocyte activation properties. Relatively preserving the native IL-8 response, as noted in the anti-CD14 treated groups, may keep intact these important mediators of the host response to LPS.

SUMMARY OF SEQUENCES

SEQ ID NO:1 is the nucleic acid and deduced amino acid sequence of the 28C5 heavy chain.

5 SEQ ID NO:2 is the deduced amino acid sequence of the 28C5 heavy chain.

SEQ ID NO:3 is the nucleic acid and deduced amino acid sequence of the 28C5 light chain.

SEQ ID NO:4 is the deduced amino acid sequence of the 28C5 light chain.

10 SEQ ID NO:5 is the nucleic acid and deduced amino acid sequence of the 18E12 heavy chain.

SEQ ID NO:6 is the deduced amino acid sequence of the 18E12 heavy chain.

SEQ ID NO:7 is the nucleic acid and deduced amino acid sequence of the 18E12 light chain.

15 SEQ ID NO:8 is the deduced amino acid sequence of the 18E12 light chain.

SEQ ID NO:9 shows the nucleic acid sequence which encodes the human soluble CD14 receptor.

SEQ ID NO:10 is the amino acid sequence of CDR1 of 3C10 heavy chain.

SEQ ID NO:11 is the amino acid sequence of CDR2 of 3C10 heavy chain.

SEQ ID NO:12 is the amino acid sequence of CDR3 of 3C10 heavy chain.

SEQ ID NO:13 is the amino acid sequence of CDR1 of 28C5 heavy chain.

SEQ ID NO:14 is the amino acid sequence of CDR2 of 28C5 heavy chain.

SEQ ID NO:15 is the amino acid sequence of CDR3 of 28C5 heavy chain.

5 SEQ ID NO:16 is the amino acid sequence of CDR1 of 3C10 light chain.

SEQ ID NO:17 is the amino acid sequence of CDR2 of 3C10 light chain.

SEQ ID NO:18 is the amino acid sequence of CDR3 of 3C10 light chain.

SEQ ID NO:19 is the amino acid sequence of CDR1 of 28C5 light chain.

SEQ ID NO:20 is the amino acid sequence of CDR2 of 28C5 light chain.

10 SEQ ID NO:21 is the amino acid sequence of CDR3 of 28C5 light chain.

SEQ ID NO:22 is the amino acid sequence of CDR1 of 23G4 light chain.

SEQ ID NO:23 is the amino acid sequence of CDR2 of 23G4 light chain.

SEQ ID NO:24 is the amino acid sequence of CDR3 of 23G4 light chain.

SEQ ID NO:25 is the amino acid sequence of the 23G4 light chain.

CLAIMS:

1. A hybridoma cell line which produces a monoclonal antibody, the monoclonal antibody being specifically reactive with cell surface CD14 and inhibiting CD14 mediated cell activation by a ligand.
2. The monoclonal antibody produced by the hybridoma cell line of claim 1.
3. The monoclonal antibody of claim 2 which has an affinity of at least about $3 \times 10^{-9} \text{M}^{-1}$.
4. The monoclonal antibody of claim 3 which is designated 28C5 or 23G4.
5. A biologically active fragment of the monoclonal antibody of claim 2 or 4.
6. The biologically active fragment of claim 5, wherein the biologically active fragment is Fab, Fab' or (Fab')₂.
7. The monoclonal antibody of claim 2, wherein the monoclonal antibody is a murine monoclonal antibody.
8. The monoclonal antibody of claim 2, wherein the monoclonal antibody is a human monoclonal antibody.
9. A chimeric antibody comprising a variable region of the monoclonal antibody of claim 2.

10. A CDR grafted antibody comprising a complementarity determining region (CDR) of the monoclonal antibody of claim 2.
11. A nucleic acid molecule encoding the monoclonal antibody of claim 2.
12. A nucleic acid molecule encoding the antibody fragment of claim 5.
13. A nucleic acid molecule encoding the chimeric antibody of claim 9.
14. A nucleic acid molecule encoding the CDR grafted antibody of claim 10.
15. The nucleic acid molecules as in any of claims 11, 12, 13, or 14, wherein the nucleic acid molecules are single stranded DNA, double stranded DNA, cDNA, or RNA.
16. The nucleic acid molecule as in any of claims 11, 12, 13, or 14 operatively linked to a promoter of RNA transcription.
17. An expression vector comprising the nucleic acid molecule of claims 11, 12, 13, or 14.
18. The expression vector of claim 17, wherein the expression vector is a virus, plasmid, or cosmid.
19. A host cell containing the expression vector of claim 17.
20. The host cell of claim 19, wherein the host cell is a procaryotic cell.
21. The host cell of claim 20, wherein the cell is a bacterial cell.

22. The host cell of claim 19, wherein the host cell is a eucaryotic cell.
23. The host cell of claim 22, wherein the eucaryotic cell is a mammalian cell or an insect cell.
24. The monoclonal antibody of claim 2, further characterized as having the ability to inhibit CD14 mediated activation of a CD14+ cell when the CD14 of the cell is bound to a ligand which induces CD14 mediated cell activation.
25. The monoclonal antibody of claim 24 which is designated 18E12.
26. A biologically active fragment of the monoclonal antibody of claims 24 or 25.
27. The biologically active fragment of claim 26, wherein the biologically active fragment is Fab, Fab' or (Fab')₂.
28. The monoclonal antibody of claim 24, wherein the monoclonal antibody is a murine monoclonal antibody.
29. The monoclonal antibody of claim 24, wherein the monoclonal antibody is a human monoclonal antibody.
30. A chimeric antibody comprising a variable region of the monoclonal antibody of claim 24.
31. A CDR grafted antibody comprising a complementarity determining region (CDR) of the monoclonal antibody of claim 24.

32. A nucleic acid molecule encoding the monoclonal antibody of claim 24.
33. A nucleic acid molecule encoding the monoclonal antibody of claim 27.
34. A nucleic acid molecule encoding the biologically active fragment of claim 26.
35. A nucleic acid molecule encoding the chimeric antibody of claim 30.
36. A nucleic acid molecule encoding the CDR grafted antibody of claim 31.
37. The nucleic acid molecule of claims 32, 33, 34, 35, or 36, wherein the nucleic acid is single stranded DNA, double stranded DNA, cDNA or RNA.
38. The nucleic acid molecule of claims 32, 33, 34, 35, or 36 operatively linked to a promoter of RNA transcription.
39. An expression vector containing the nucleic acid molecule of claims 32, 33, 34, 35, or 36.
40. The expression vector of claim 39, wherein the expression vector is a virus, plasmid, or cosmid.
41. A host cell containing the expression vector of claim 39.

42. The host cell of claim 41, wherein the host cell is a procaryotic cell.
43. The host cell of claim 42, wherein the cell is a bacterial cell.
44. The host cell of claim 41, wherein the host cell is a eucaryotic cell.
45. The host cell of claim 44, wherein the cell is a mammalian cell or an insect cell.
46. A pharmaceutical composition comprising the antibody of claim 2 in a pharmaceutically acceptable carrier.
47. A method of detecting CD14 in a sample which comprises contacting the sample with an antibody or fragment thereof which is specifically reactive with cell surface CD14 and soluble CD14 and determining whether the antibody binds to the sample.
48. The method of claim 47, wherein the antibody has the specificity of 28C5, 23G4, or 18E12.
49. The method of claim 48, wherein the antibody is detectably labeled.
50. The method of claim 49, wherein the detectable label is selected from the group consisting of a radioisotope, a fluorescent compound, a colloidal metal, a chemiluminescent compound, a bioluminescent compound and an enzyme.
51. The method of claim 47, wherein the method of detecting is by a competitive immunoassay method.

52. The method of claim 47, wherein the sample comprises a body fluid.
53. A method of inhibiting the binding of LPS to CD14 which comprises contacting the CD14 with the monoclonal antibody of claim 4 or a biologically active fragment thereof.
54. A method of inhibiting the binding of an LPS/CD14 complex to a cell, which comprises contacting the complex with the monoclonal antibody of claim 2 or a biologically active fragment thereof.
55. A method of inhibiting CD14 mediated activation of a cell expressing CD14 receptor, which comprises contacting the cell with an effective amount of the antibody of claim 2 or a biologically active fragment thereof.
56. The method of claim 55, wherein the antibody allows binding of the ligand to CD14.
57. The method of claim 56, wherein the antibody allows at least about 50% binding of the ligand to CD14.
58. The method of claim 56, wherein the antibody allows at least about 80% binding of the ligand to CD14.
59. The method of claim 56, wherein the antibody has the specificity of monoclonal antibody 18E12.
60. The method of claim 55, wherein the antibody has the specificity of monoclonal antibody 28C5 or 23G4.

61. The method of claim 55, wherein the CD14 mediated activation is associated with NF- κ B activation.
62. The method of claim 61, wherein the NF- κ B activation is associated with sepsis.
63. The method of claim 55, wherein the ligand is LPS.
64. The method of claim 55, wherein the cell is an animal or a human patient cell.
65. The method of claim 55, wherein an effective amount is from about .25 mg/kg/body weight to about 50 mg/kg/body weight.

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+1 ATGGAGCGCG CGTCCTGCTT GTTGCTGCTG CTGCTGCCGC TGGTGCACGT
+51 CTCTGCGACC ACGCCAGAAC CTTGTGAGCT GGACGATGAA GATTTCCGCT
+101 GCGTCTGCAA CTTCTCCGAA CCTCAGCCCG ACTGGTCCGA AGCCTTCCAG
+151 TGTGTGTCTG CAGTAGAGGT GGAGATCCAT GCCGGCGGTC TCAACCTAGA
+201 GCCGTTTCTA AAGCGCGTCG ATGCGGACCG CGACCCGCGG CAGTATGCTG
+251 ACACGGTCAA GGCTCTCCGC GTGCGGCGGC TCACAGTGGG AGCCGCACAG
+301 GTTCCTGCTC AGCTACTGGT AGGCGCCCTG CGTGTGCTAG CGTACTCCCG
+351 CCTCAAGGAA CTGACGCTCG AGGACCTAAA GATAACCGGC ACCATGCCTC
+401 CGCTGCCTCT GGAAGCCACA GGACTTGAC TTTCCAGCTT GCGCCTACGC
+451 AACGTGTCGT GGGCGACAGG GCGTTCTTGG CTCGCCGAGC TGCAGCAGTG
+501 GCTCAAGCCA GGCCTCAAGG TACTGAGCAT TGCCCAAGCA CACTCGCCTG
+551 CCTTTTCCTG CGAACAGGTT CGCGCCTTCC CGGCCCTTAC CAGCCTAGAC
+601 CTGTCTGACA ATCCTGGACT GGGCGAACGC GGACTGATGG CGGCTCTCTG
+651 TCCCCACAAG TTCCCGGCCA TCCAGAATCT AGCGCTGCGC AACACAGGAA
+701 TGGAGACGCC CACAGGCGTG TGCGCCGCAC TGGCGGCGGC AGGTGTGCAG
+751 CCCCACAGCC TAGACCTCAG CCACAACCTCG CTGCGCGCCA CCGTAAACCC
+801 TAGCGCTCCG AGATGCATGT GGTCCAGCGC CCTGAACTCC CTCAATCTGT
+851 CGTTCGCTGG GCTGGAACAG GTGCCTAAAG GACTGCCAGC CAAGCTCAGA
+901 GTGCTCGATC TCAGCTGCAA CAGACTGAAC AGGGCGCCGC AGCCTGACGA
+951 GCTGCCCCGAG GTGGATAACC TGACACTGGA CGGGAATCCC TTCCTGGTCC
+1001 CTGGAAGTGC CCTCCCCAC GAGGGCTCAA TGAAGTCCGG CGTGGTCCCA
+1051 GCCTGTGCAC GTTCGACCCT GTCGGTGGGG GTGTCGGGAA CCCTGGTGCT
+1101 GCTCCAAGGG GCCCGGGGCT TTGCCTAA

FIG. 1

5' ⁴ CCC CTC GAG CTT CAG CAG TCA GGA CCT GGC CTG GTG AAA CCT TCT CAG TCT CTG TCC CTC ACC TGC ACT GTC
 pro pro leu glu leu gln gln ser gly pro gly leu val lys pro ser gln ser leu ser leu thr cys thr val
 ACT GGC TAC TCA ATC ACC AGT GAT TCT GCC TGG AAC TGG ATC ^{35B} Fr.2 CDR ³¹ thr gly tyr ser ile thr ser asp ser ala trp asn trp ile arg gln phe pro gly asn arg leu glu trp
⁵⁰ ATG GGC TAC ATA AGC TAC AGT GGT AGC ACT AGC TAC AAC CCA TCT CTC AAA AGT ⁶⁵ Fr.3 ³ met gly tyr ile ser tyr ser gly ser thr ser leu lys ser arg ile ser ile
 ACT CGA GAC ACA TCC AAG AAC CAG TTC TTC CTG CAG TTG AAT TCG GTG ACT ACT GAG GAC ACA GCC ACA TAT TAC
 thr arg asp thr ser lys asn gln phe phe leu gln leu asn ser val thr thr glu asp thr ala thr tyr tyr
⁹⁵ TGT GTA AGA GGG CTC CGG TTT GCT TAC TGG GGG AAG GGG ACT CTG GTC ACT GTC TCT GCA GCA AAA ACA ACC
 cys val arg gly leu arg phe ala tyr trp gly gln gly thr leu val thr val ser ala ala lys thr thr ¹¹³ CH-1 ^{2/28}
 CCC CCC TCT GTC TAT CCA CTG CCC CCT GGA TCT GCT GCC CAA ACT AAC TCC ATG GTG ACC CTG GGA TGC CTG GTC
 pro pro ser val tyr pro leu pro pro gly ser ala ala gln thr asn ser met val thr leu gly cys leu val
 AAG GCC TAT TTC CCT GAG CCA GTG ACA GTG ACC TGG AAC TCT GGA TCC CTG TCC AGC GGT TGG CAC ACC TTC CCA
 lys gly tyr phe pro glu pro val thr val thr trp asn ser gly ser leu ser ser gly val his thr phe pro
 GCT GTC CTG CAG TCT GAC CTC TAC ACT CTG AGC TCA GTG ACT GTC CCC TCC AGC ACC TGG CCC AGC GAG ACC
 ala val leu gln ser asp leu tyr thr leu ser ser val thr val pro ser thr trp pro ser glu thr
 GTC ACC TGC AAC GTT GCC CAC CCG GCC AGC AGC ACC AAG GTG GAC AAG AAA ATT 3' ²²³
 val thr cys asn val ala his pro ala ser thr lys val asp lys lys ile

FIG. 2

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5' 5 CAG TCT CCA GCT TCT TTG GCT GTG TCT CTA GGG CAG AGG GCC CCA TAT CCT TGC AGA GCC AGT GAA CDR1
 met thr gln ser pro ala ser leu ala val ser leu gly gln arg ala thr ile ser cys arg ala ser glu

AGT GTT GAT AGT TAT GTC AAT AGT TTT CTC CAC TGG TAC CAG CAG AAA CCA GGA CAG CCA CCC AAA CTC
 ser val asp ser tyr val asn ser phe leu his trp tyr gln gln lys pro gly gln pro pro lys leu Fr.2

CTC ATC TAT CGT GCA TCC AAC CTA CAA TCT GGG ATC CCT GCC AGG TTC AGT GGC AGT GGG TCT AGG ACA GAC
 leu ile tyr arg ala ser asn leu gln ser gly ile pro ala arg phe ser gly ser gly ser arg thr asp Fr.3

TTC ACC CTC ACC ATT AAT CCT GTG GAG GCT GAT GAT GTT GCA ACC TAT TAC TGT CAG CAA AGT AAT GAG GAT
 phe thr leu thr ile asn pro val glu ala asp asp val ala thr tyr cys gln gln ser asn glu asp CDR3

CCG ACG ACG TCG GGA GGC ACC AAG CTG GAA ATA AAA CGG GCT GAT GCT GCA CCC CTT GTA TCC ATC TTC CCC
 pro thr thr phe gly gly thr lys leu glu ile lys arg ala asp ala ala pro leu val ser ile phe pro 107 CH-1

CCA TCC AGT GAG CAG TTA ACA TCT GGA GGT GCC TCA GTT GTG TGC TTC TTG AAC AAC TTC TAC CCC AAA GAC ATC
 pro ser ser glu gln leu thr ser gly gly ala ser val val cys phe leu asn asn phe tyr pro lys asp ile Fr.4

AAT GTC AAG TGG AAG ATT GAT GTC AGT GAA CGA CAA AAT GGC GTC CTG AAC AGT TGG ACT GAT CAG GAC AGC AAA
 asn val lys trp lys ile asp val ser glu arg gln asn asn gly val leu asn ser trp thr asp gln asp ser lys

GAC AGC ACC TAC AGC ATG AGC ACC CTC ACG TTG ACC AAG GAC GAG TAT GAA CGA CAT AAC AGC TAT ACC TGT
 asp ser thr tyr ser met ser thr leu thr leu thr lys asp glu tyr glu arg his asn ser tyr thr cys

GAG GCC ACT CAC AAG ACA TCA ACT TCA CCC ATT GTC AAG AGC TTC AAC AGG AAT GAG TGT 3'
 glu ala thr his lys thr ser thr ser pro ile val lys ser phe asn arg asn glu cys 214

FIG. 3

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5' a.a.5 Fr. 1
 Pro Pro Leu Glu Ser Gly Pro Gly Leu Val Ala Pro Ser Gln Ser Leu Ser Ile Thr Cys Thr Val
 CCC CCC CTC GAG TCA GGA CCT GGC CTG GCG GCG GCG CCC TCA CAG AGC CTG TCC ATT ACC TGC ACT GTC

 Ser Gly Phe Ser Leu Thr Asn Tyr Asp Ile Ser Trp Ile Arg Gln Pro Pro Gly Lys Gly Leu
 TCT GGG TTC TCA TTA ACC AAC TAT GAT ATA AGC TGG ATT CGC CAG CCA CCA GGA AAA GGT CTG
 31 CDR1 36 Fr. 2

 Glu Trp Leu Gly Val Ile Trp Thr Ser Gly Gly Thr Asn Tyr Asn Ser Ala Phe Met
 GAG TGG CTT GGA GTA ATA TGG ACT AGT GGA GGC ACA AAT TAT AAT TCA GCT TTC ATG
 50 CDR2

 Ser Arg Leu Ser Ile Thr Lys Asp Asn Ser Lys Ser Gln Val Phe Leu Lys Met Asn Gly Leu Gln
 TCC CGA CTG AGC ATC ACC ACC AAG GAC AAC TCC AAG AGC CAA GTT TTC TTA AAA ATG AAC GGT CTG CAA
 66 Fr. 3

 Thr Asp Asp Thr Gly Ile Tyr Tyr Cys Val Arg Gly Asp Gly Asn Phe Tyr Leu Tyr Asn
 ACT GAT GAC ACA GGC ATA TCT TCT TAC TGT GTA AGA GGT GAT GGT AAC TTC TAC TTG TAC AAC
 95 CDR3

 Phe Asp Tyr Trp Gly Gln Gly Thr Thr Leu Thr Thr Val Ser Ser Ala Lys Thr Thr Pro Pro Ser
 TTT GAC TAT TGG GGC CAA GGC ACC ACT CTC ACT CTC ACA GTC TCC TCA GCC AAA ACG ACA CCC CCA TCT
 103 Fr. 4 CH-1

 GTC TAT CCA CTG GCC CCT GGA TCT GCT GCC CAA ACT AAC TCC ATG GTG ACC CTG GGA TGC CTG GTC
 Val Tyr Pro Leu Ala Pro Gly Ser Ala Ala Gln Thr Asn Ser Met Val Thr Leu Gly Cys Leu Val

 AAG GGC TAT TTC CCT GAG CCA GTG ACA GTG ACC TGG AAC TCT GGA TCC CTG TCC AGC GGT GTG CAC
 Lys Gly Tyr Phe Pro Glu Pro Val Thr Val Thr Trp Asn Ser Gly Ser Leu Ser Ser Ser Ser Ser

 ACC TTC CCA GCT GTC CTG CAG TCT GAC CTC TAC ACT CTG AGC AGC TCA GTG ACT GTC CCC TCC AGC
 Thr Phe Pro Ala Val Leu Gln Ser Asp Leu Tyr Thr Leu Ser Ser Val Thr Val Thr Val Pro Ser Ser

 ACC TGG CCC AGC GAG ACC GTC ACC TGC AAC GTT GCC CAC CCG GCC AGC AGC ACC AAG GTG GAC AAG
 Thr Trp Pro Ser Glu Thr Val Thr Cys Asn Val Ala His Pro Ala Ser Ser Thr Lys Val Asp Lys

 AAA ATT 3'
 Lys Ile

FIG. 4

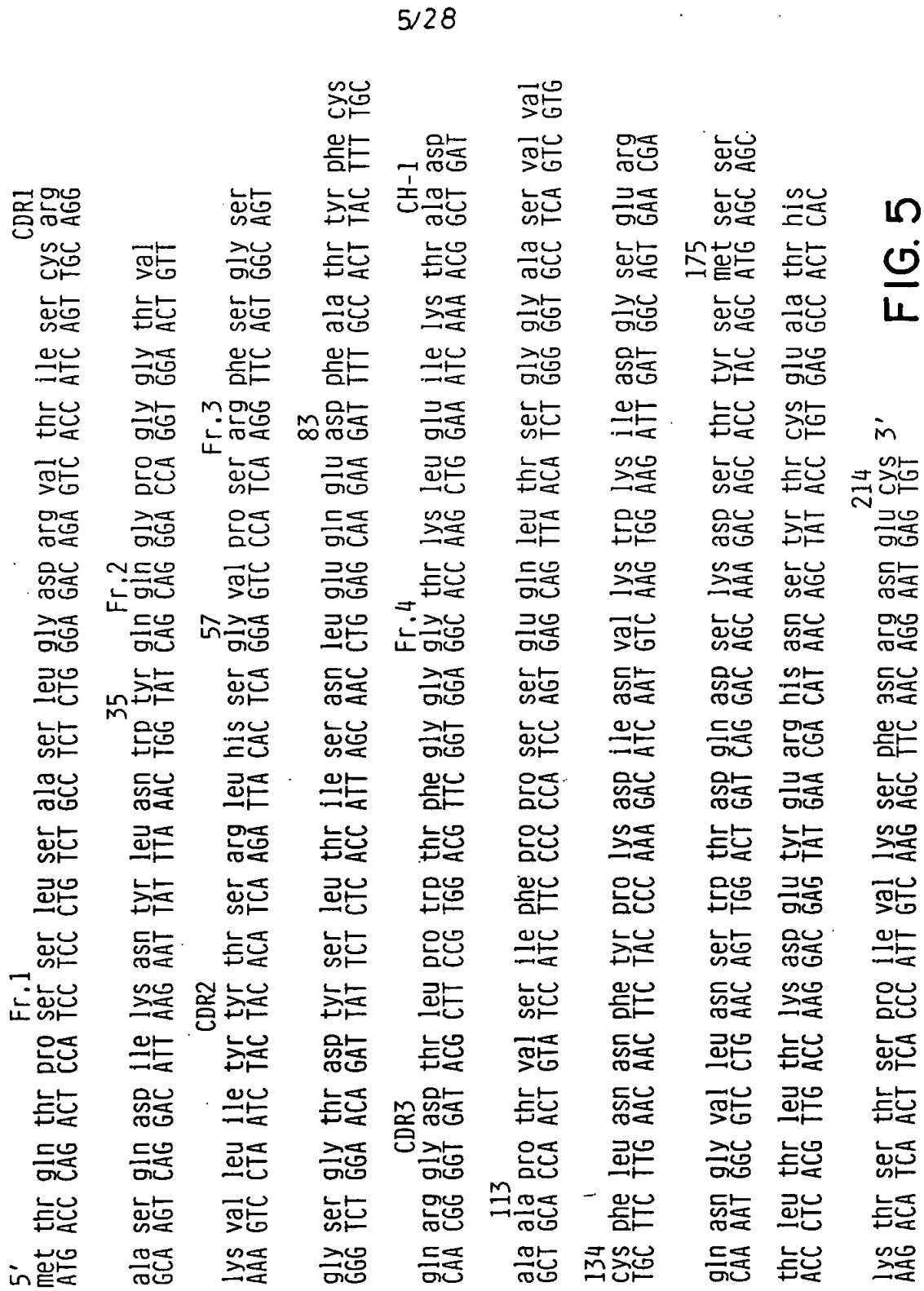


FIG. 5

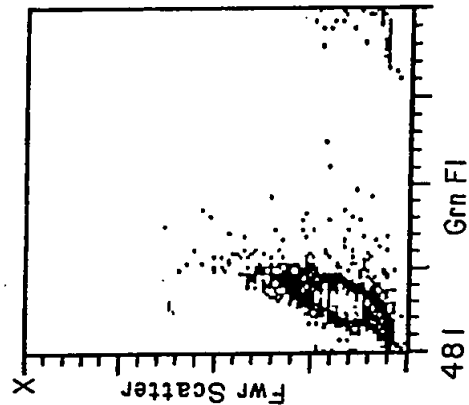


FIG. 6B

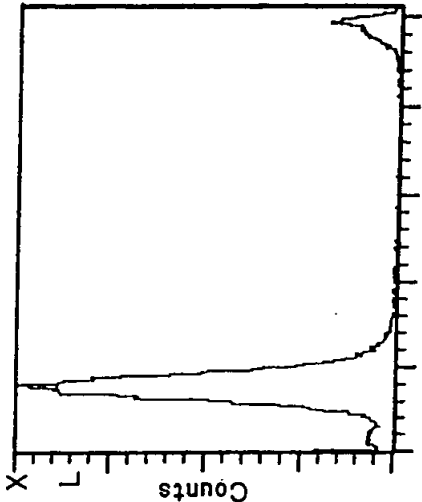


FIG. 6D

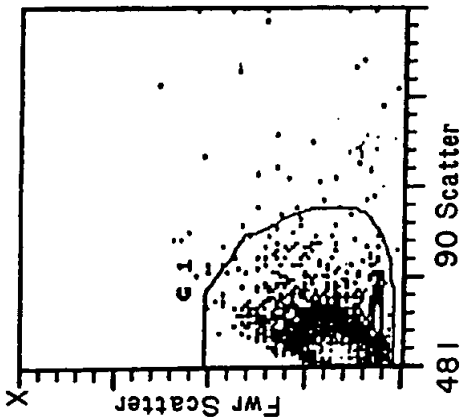


FIG. 6A

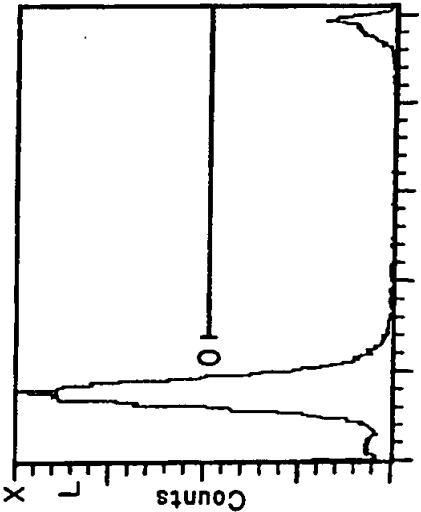


FIG. 6C

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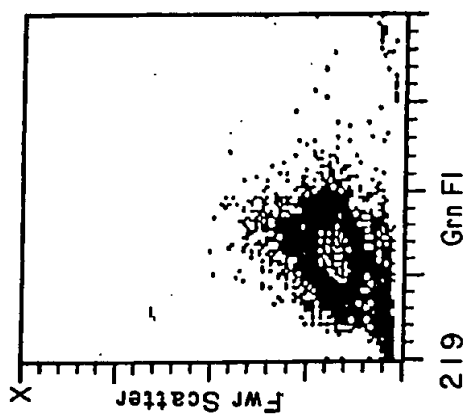


FIG. 7A

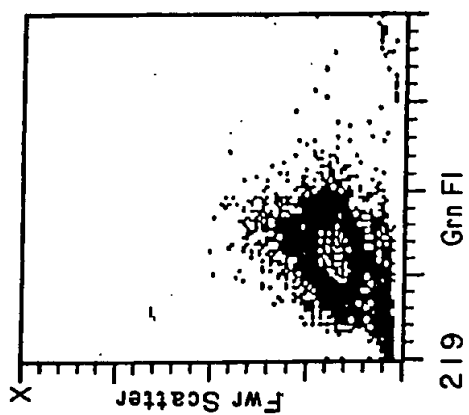


FIG. 7B

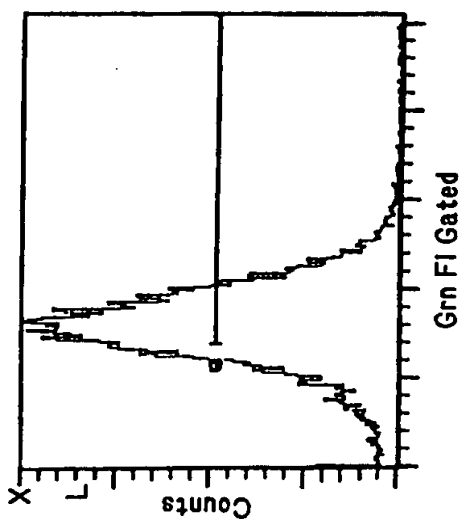


FIG. 7C

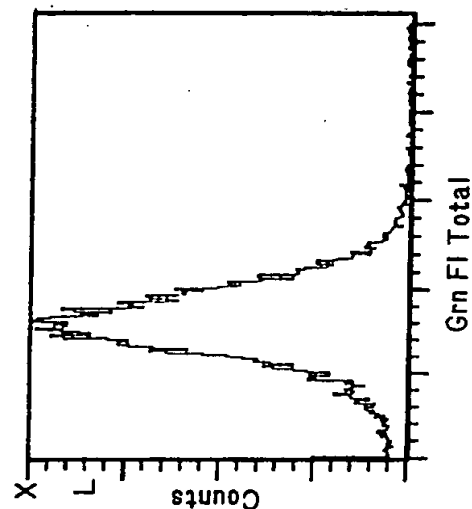


FIG. 7D

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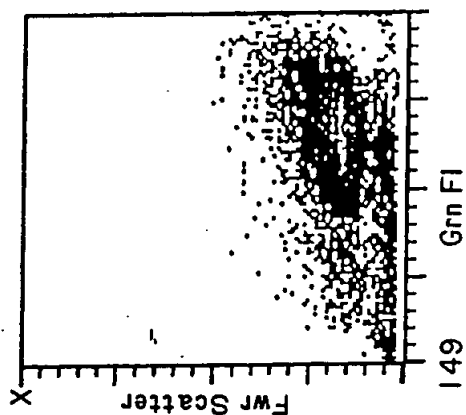


FIG. 8B

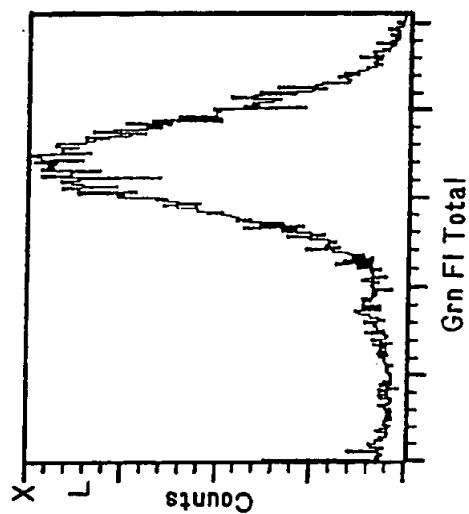


FIG. 8D

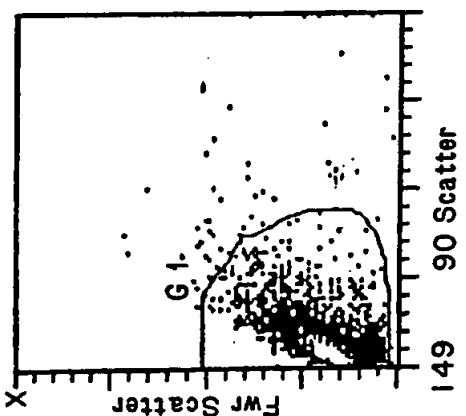


FIG. 8A

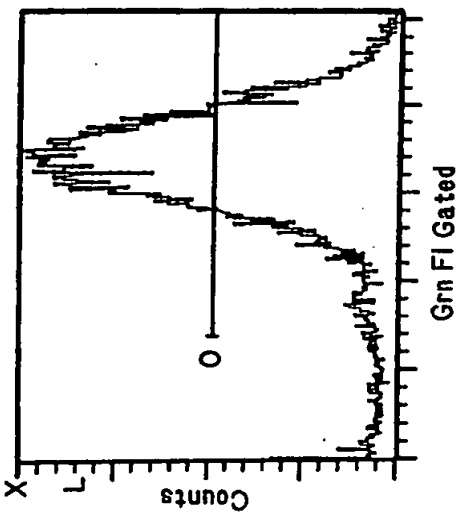


FIG. 8C

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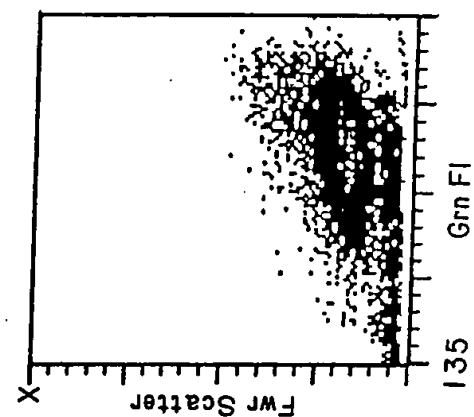


FIG. 9B

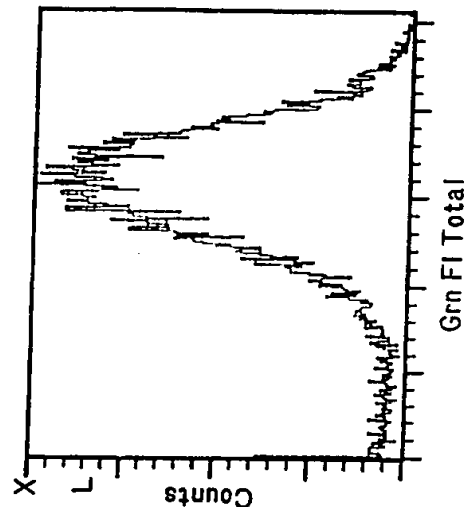


FIG. 9D

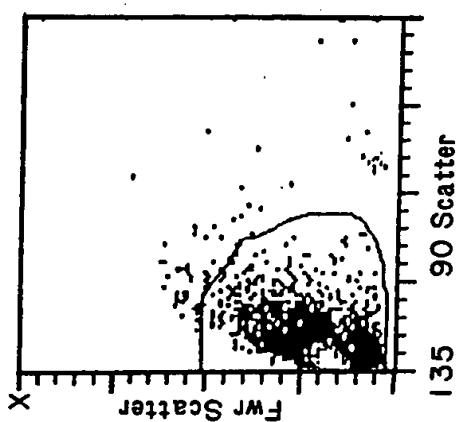


FIG. 9A

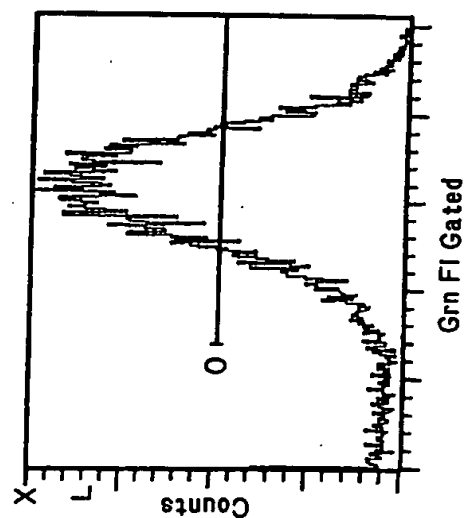


FIG. 9C

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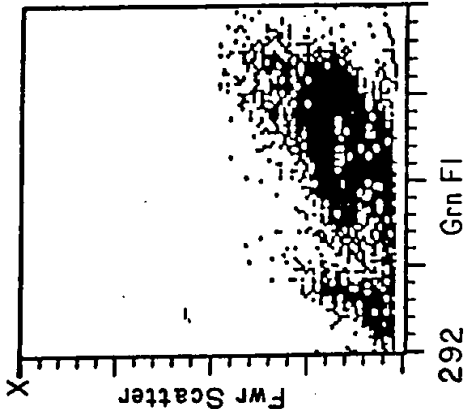


FIG. 10B

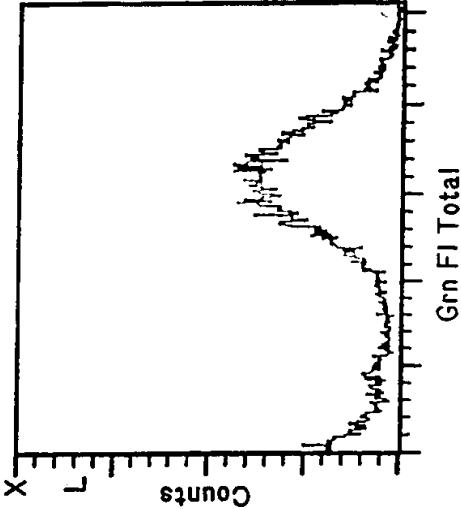


FIG. 10D

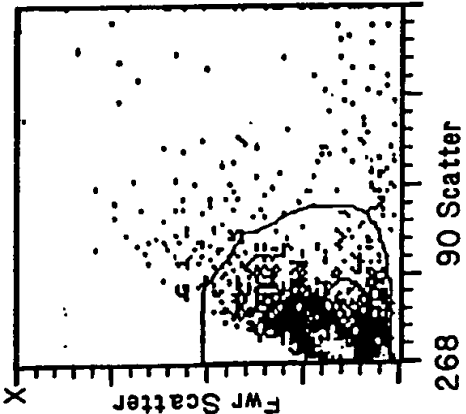


FIG. 10A

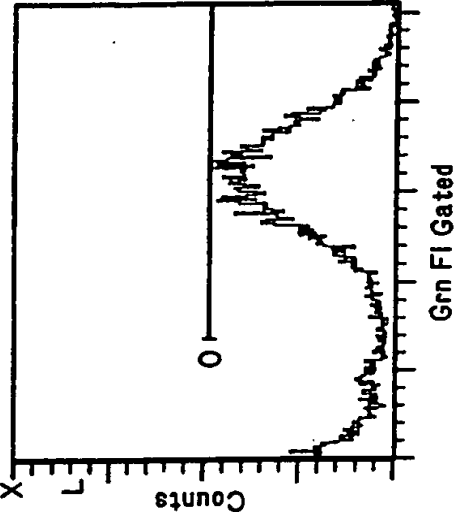


FIG. 10C

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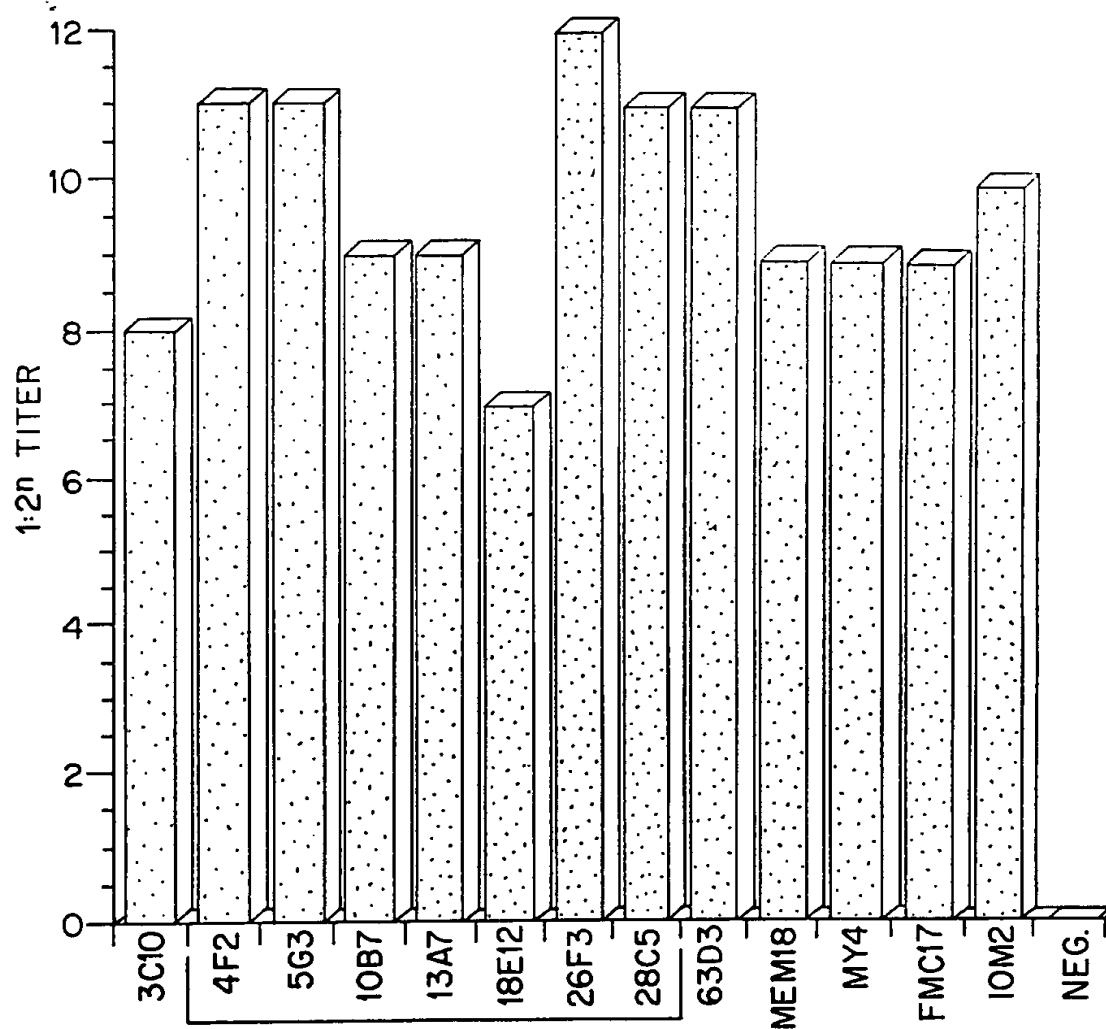


FIG. 11

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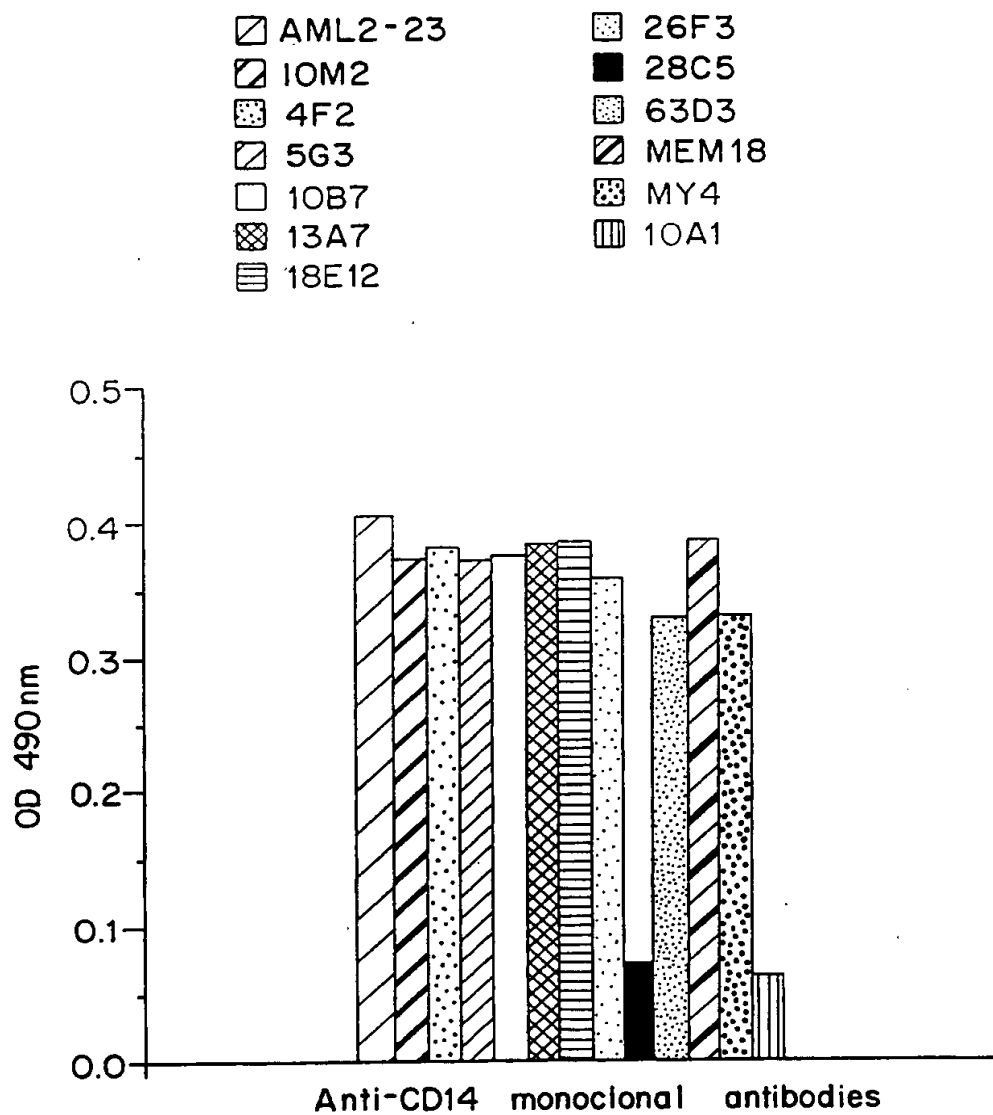


FIG. 12

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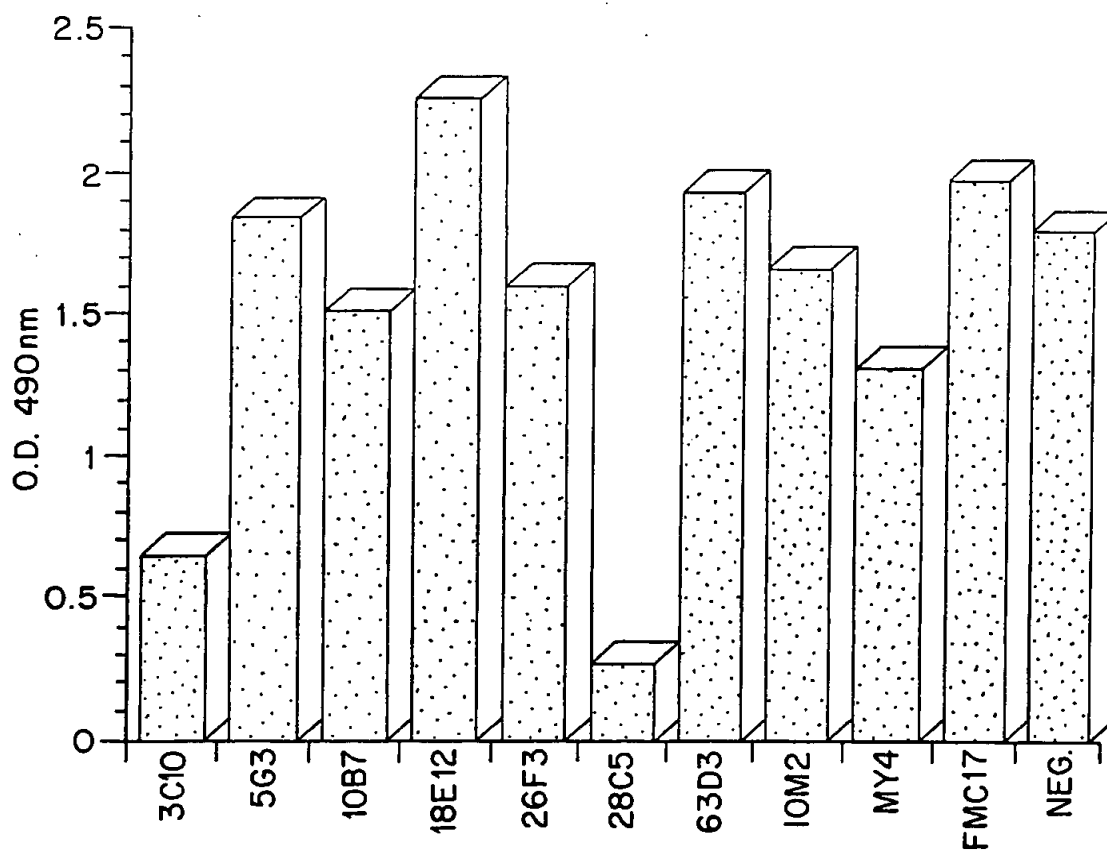


FIG. 13

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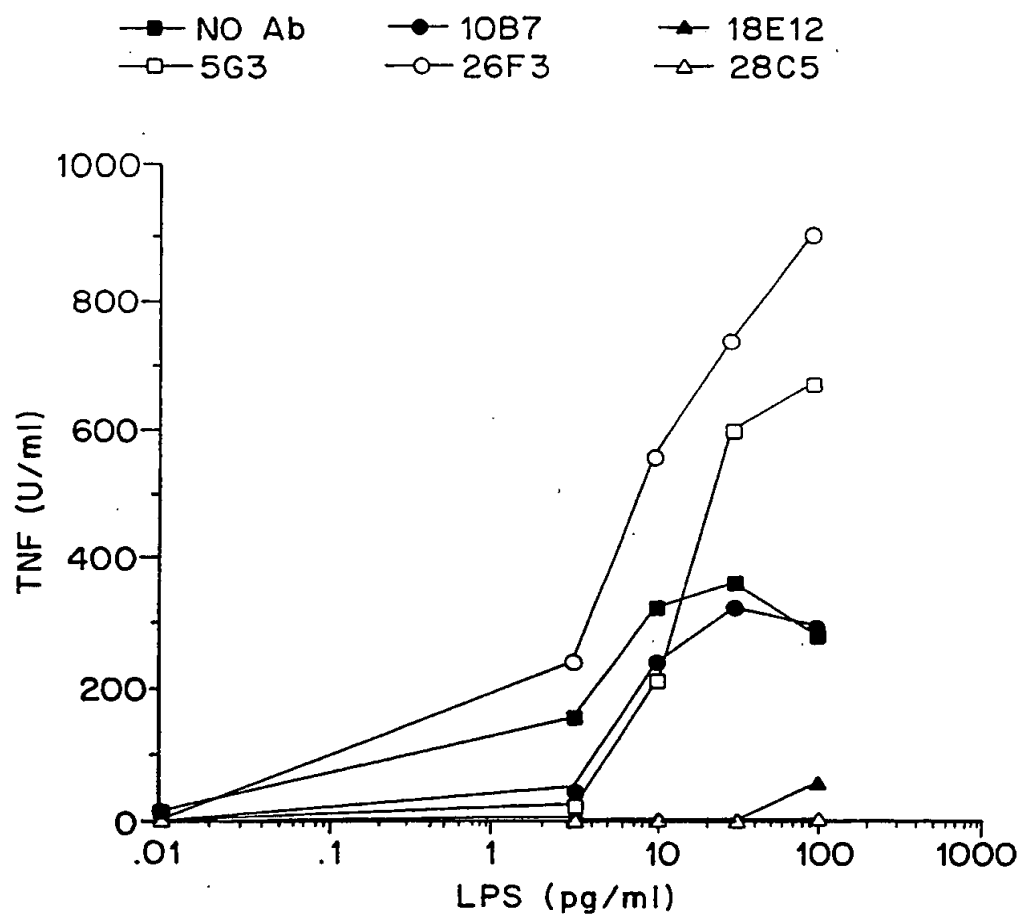


FIG. 14

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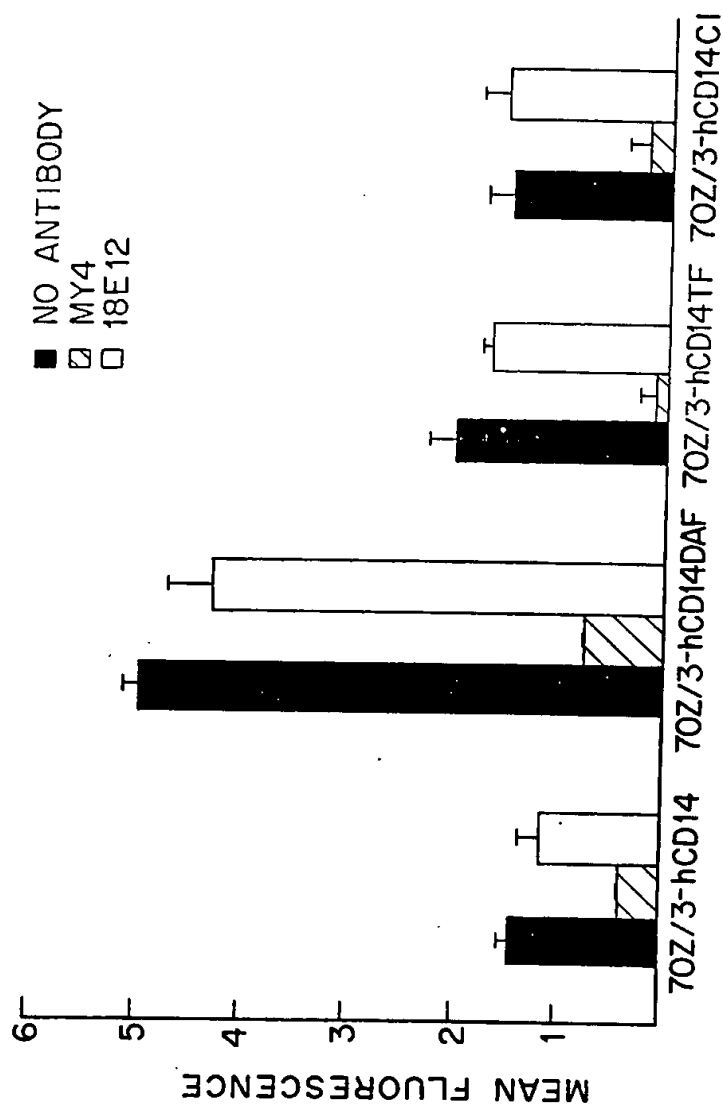


FIG. 15

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MY4	-	+	-	-	+	-	-	+	-
18E12	-	-	+	-	-	+	-	-	+
TAXOL	-	-	-	+	+	+	-	-	-
LPS	-	-	-	-	-	-	+	+	+

70Z/3-RSV

70Z/3-hCD14

70Z/3-hCD14DAF

70Z/3-hCD14TF

70Z/3-hCD14CI

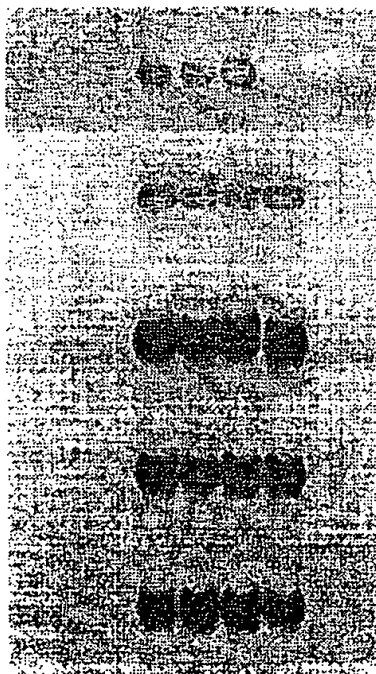
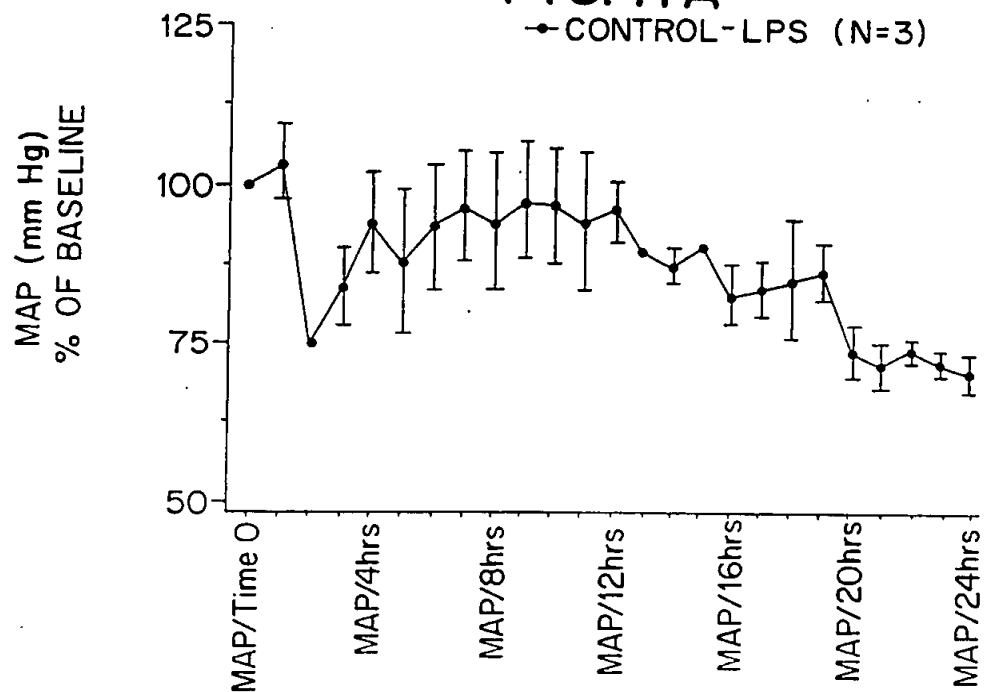
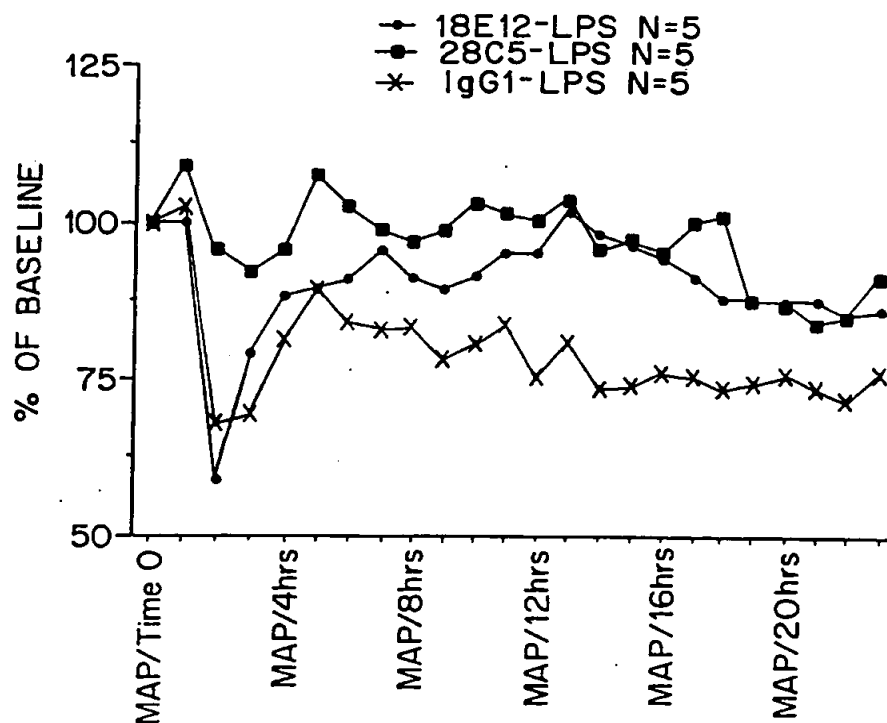


FIG. 16

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FIG. 17A**FIG. 17B**

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FIG. 18A

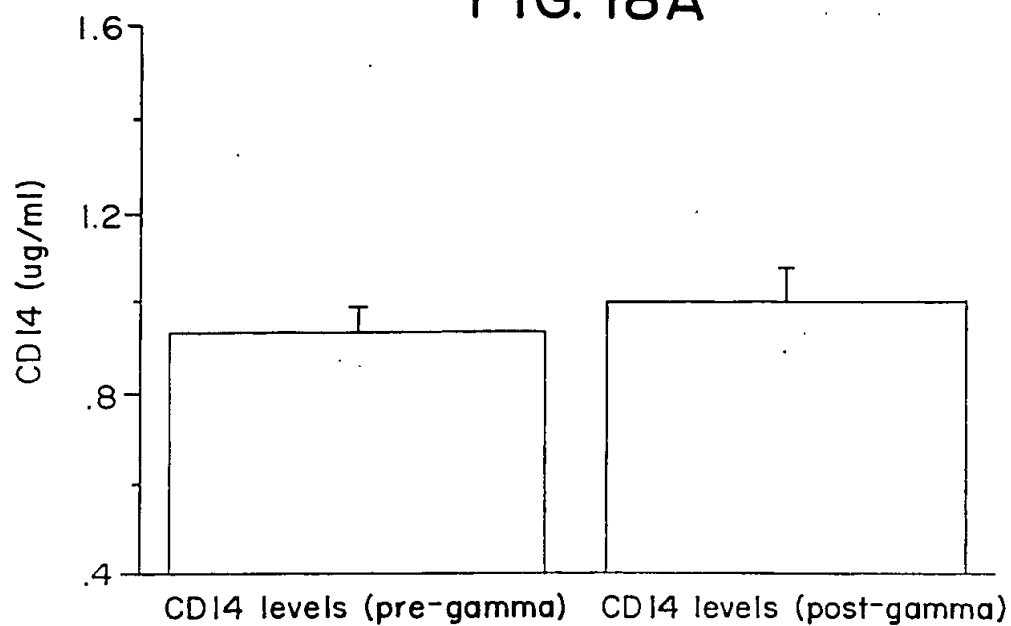
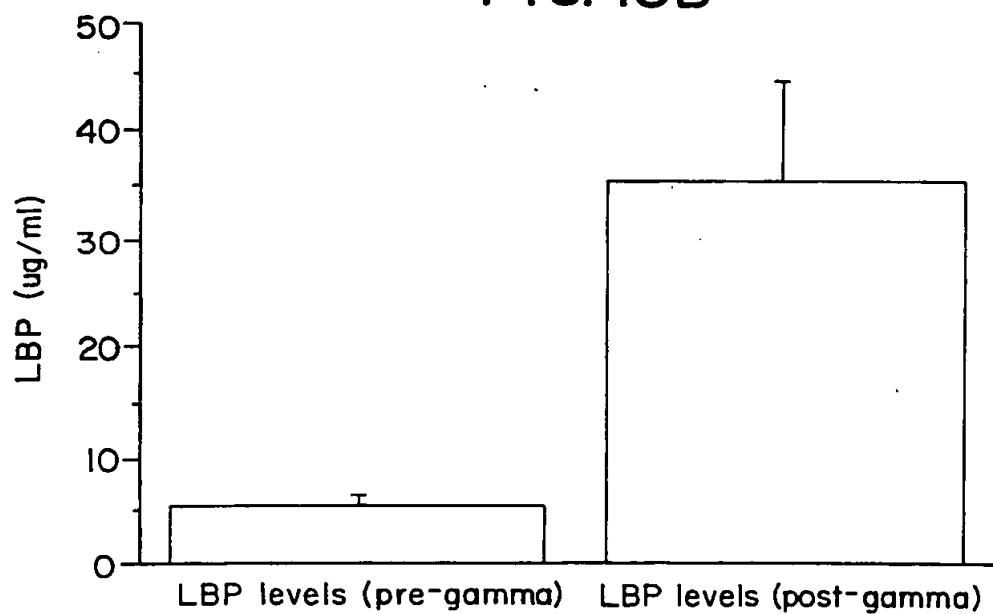


FIG. 18B



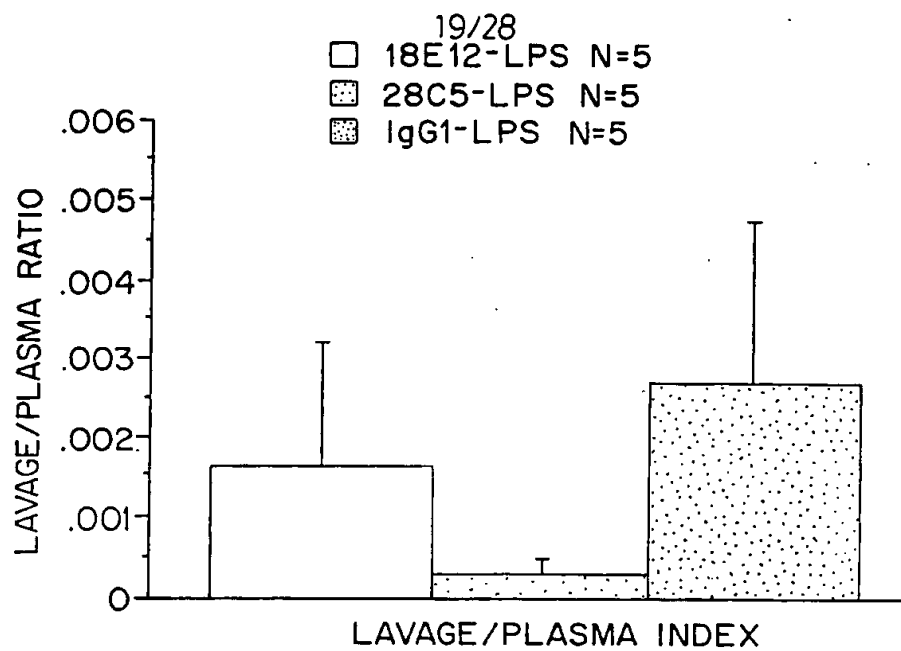


FIG. 19

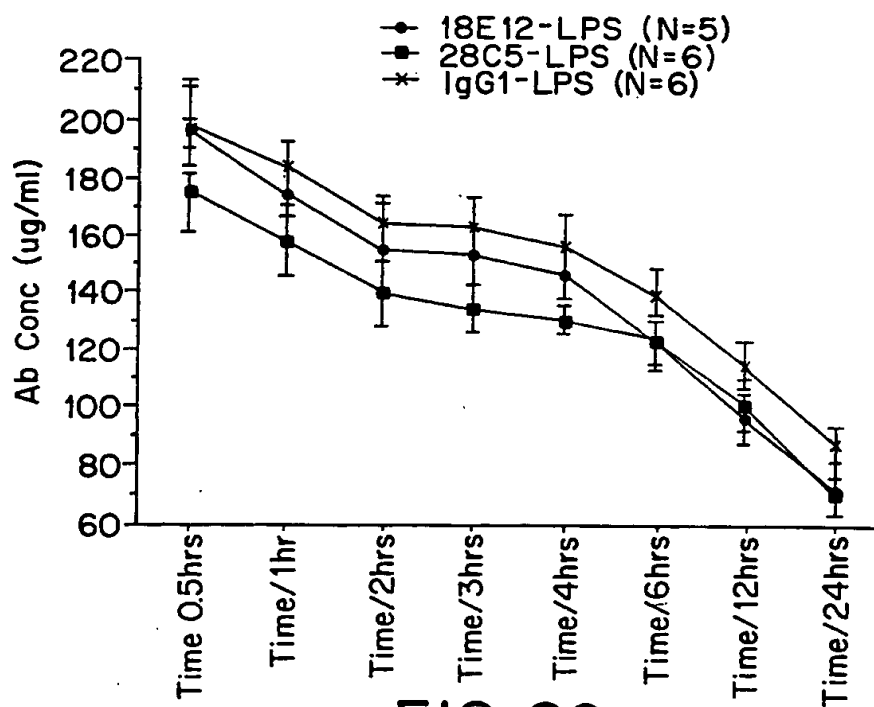


FIG. 20

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FIG. 21A

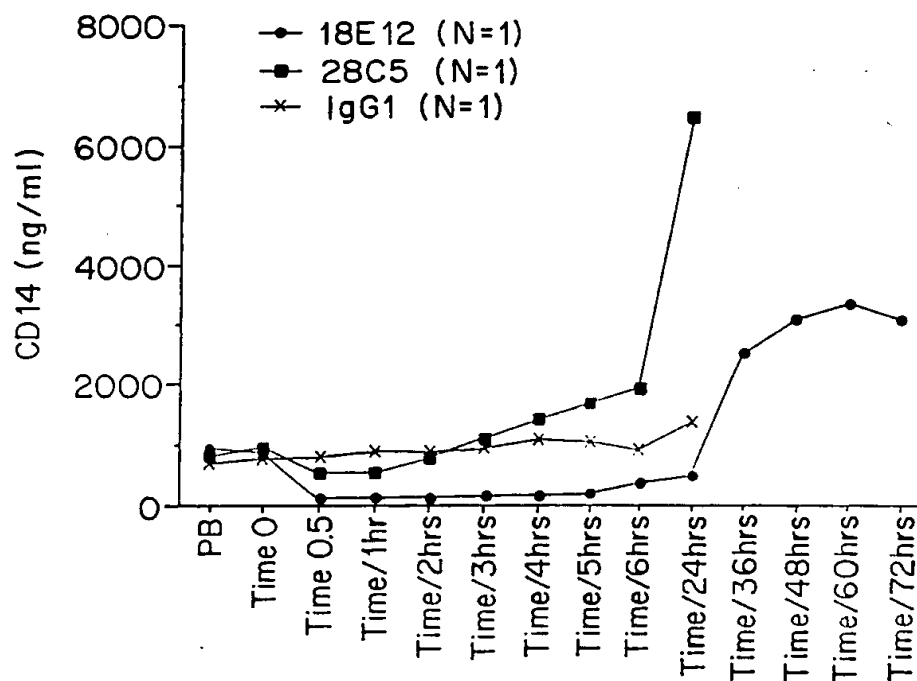
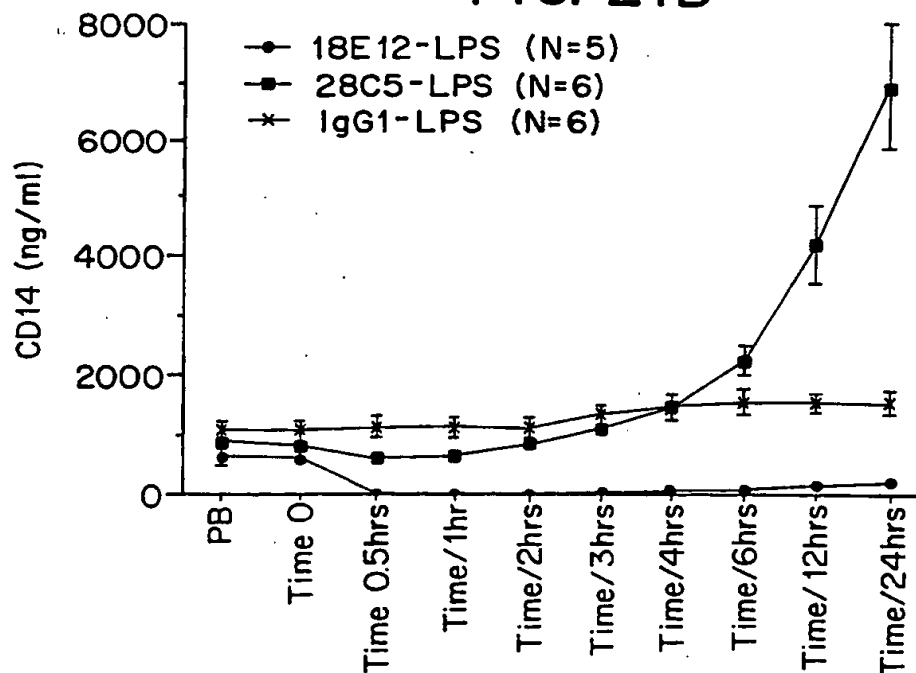


FIG. 21B



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FIG. 22A

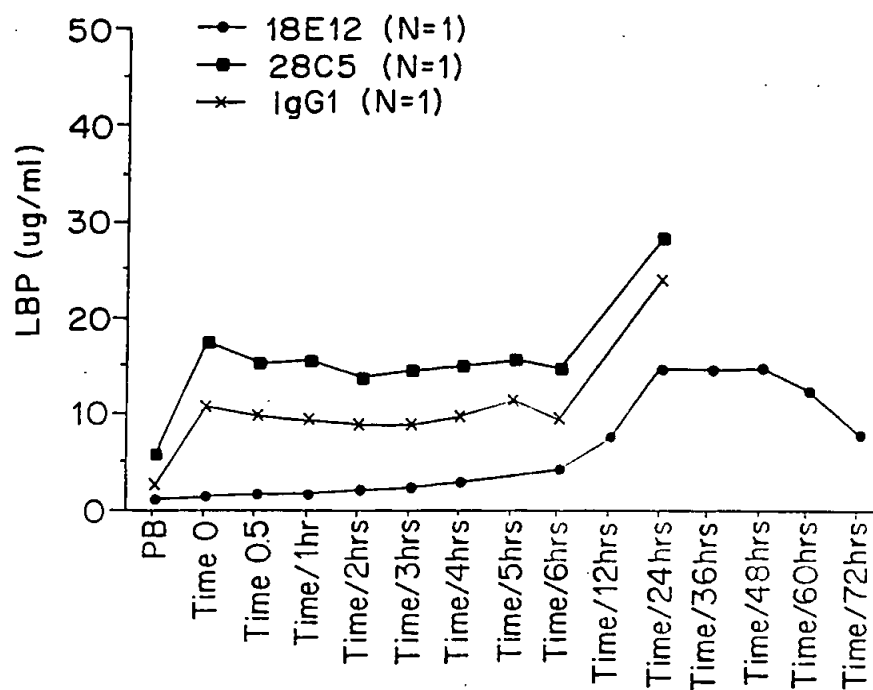
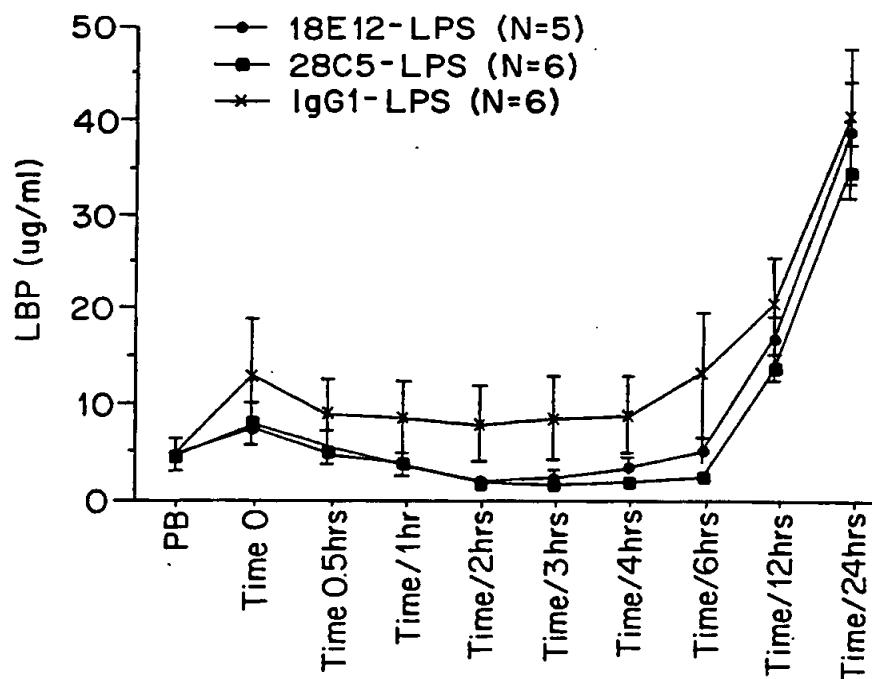


FIG. 22B



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FIG. 23A

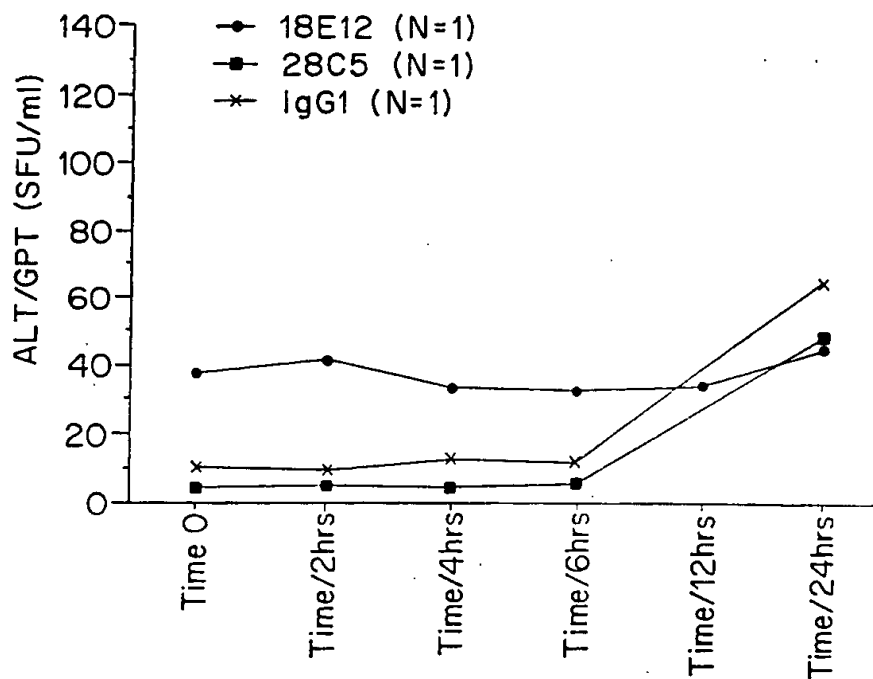
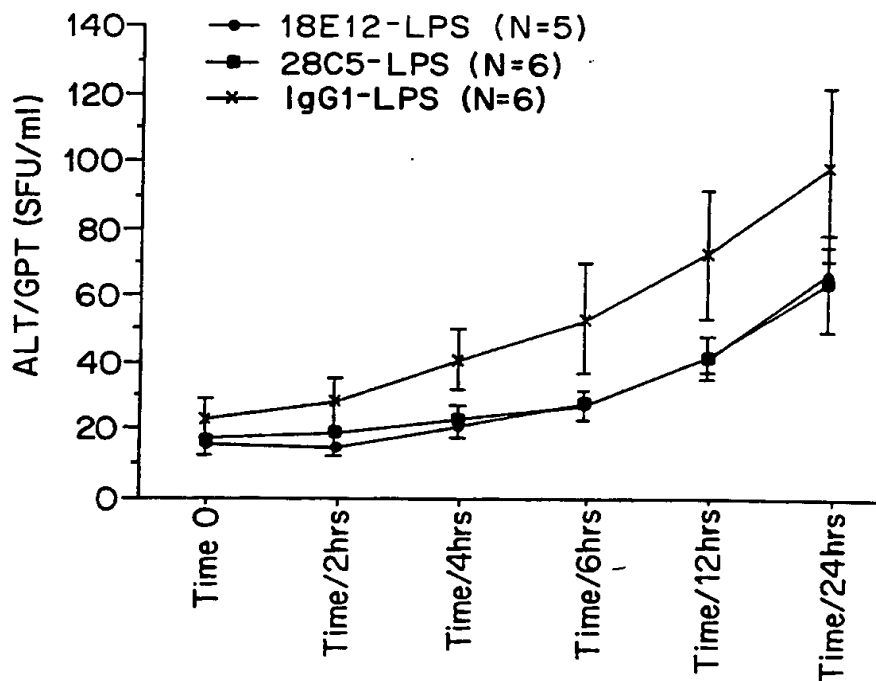


FIG. 23B



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FIG. 24A

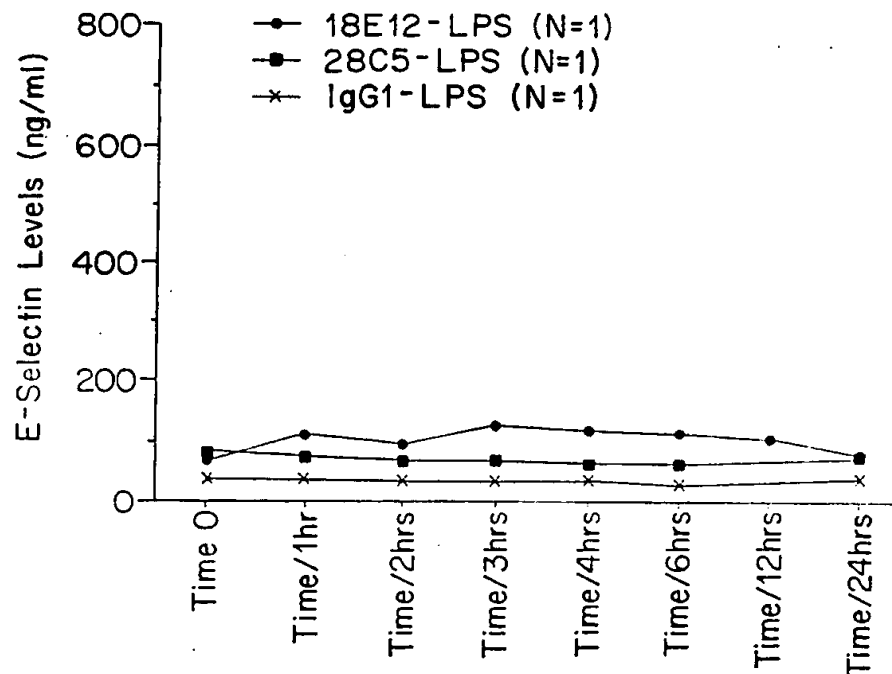
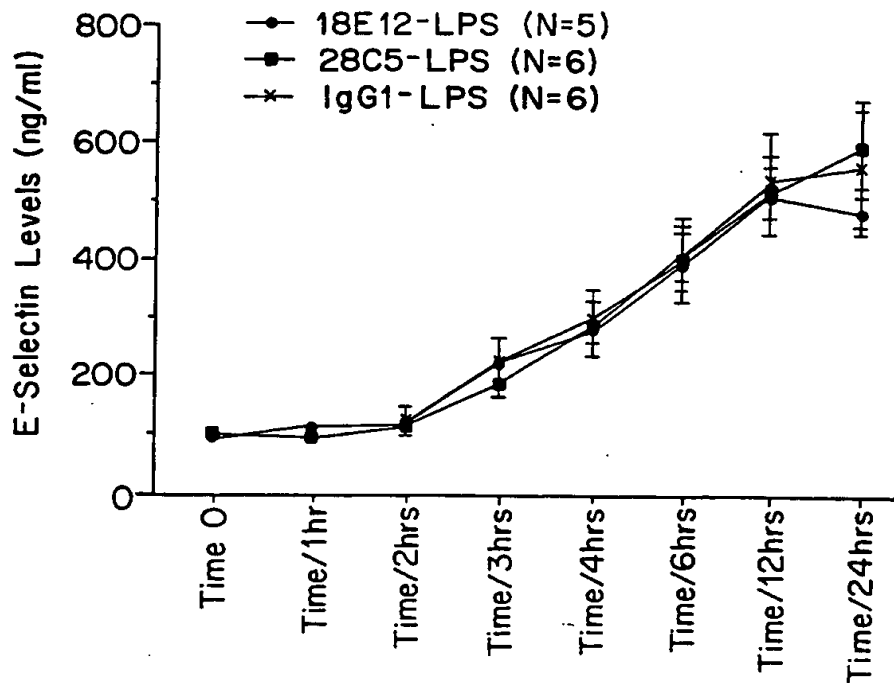


FIG. 24B



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FIG. 25A

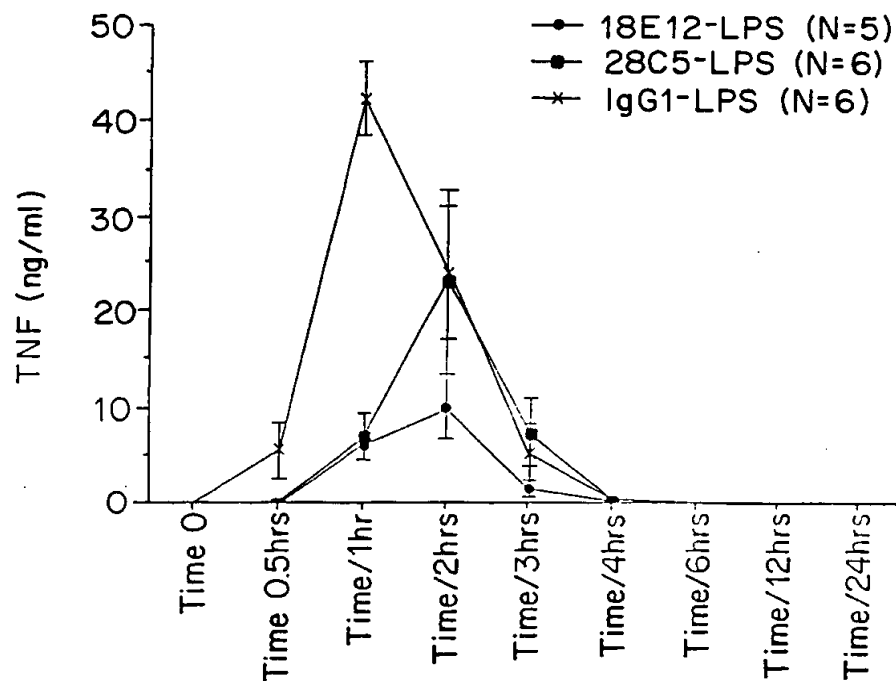
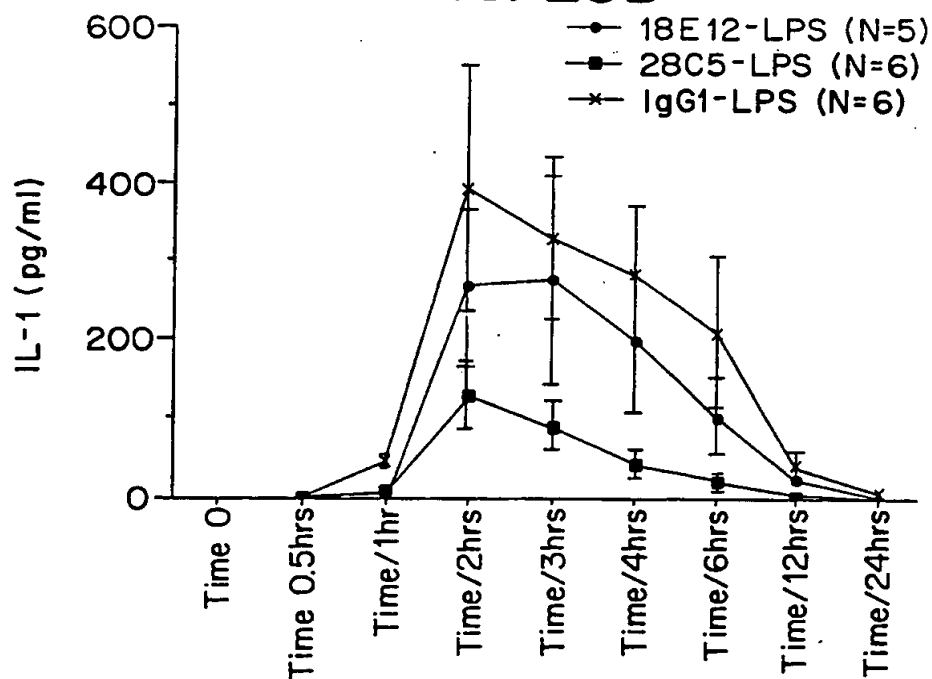


FIG. 25B



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FIG. 26A

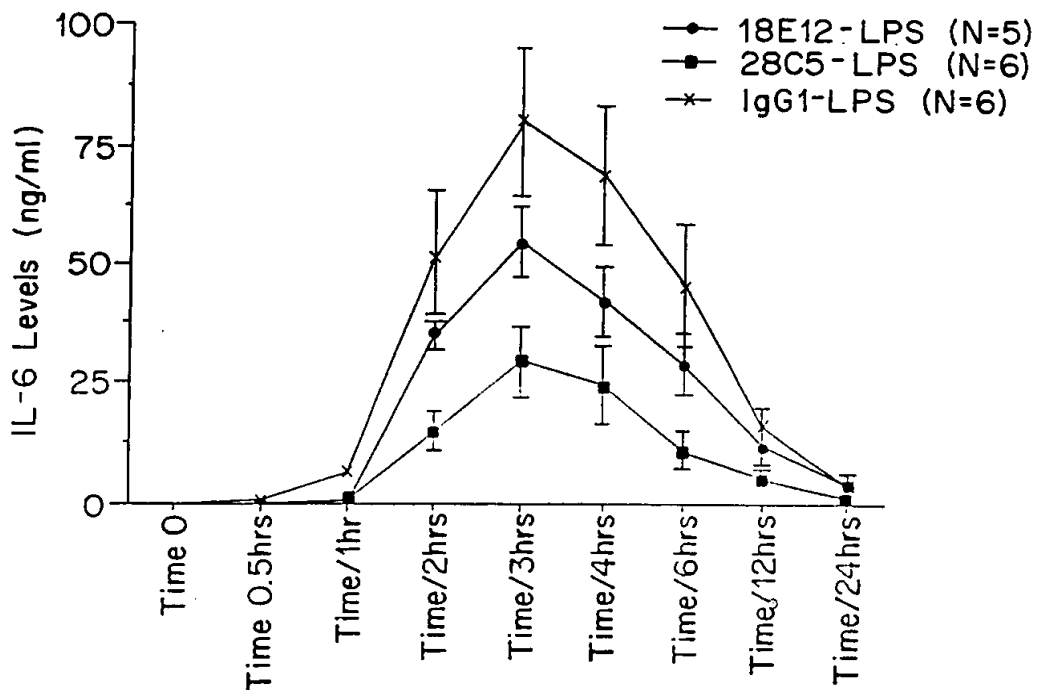
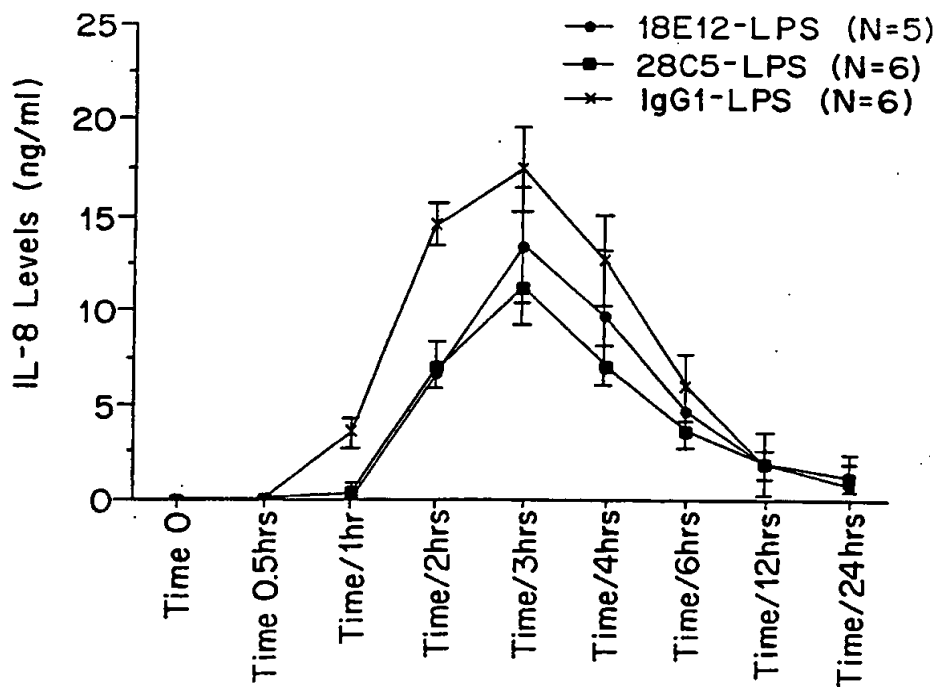


FIG. 26B



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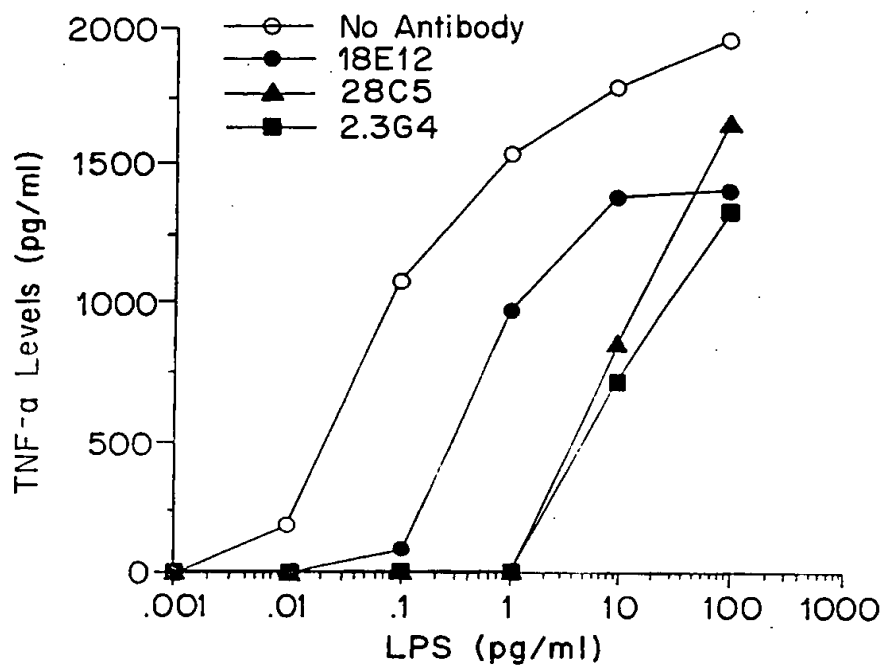


FIG. 27

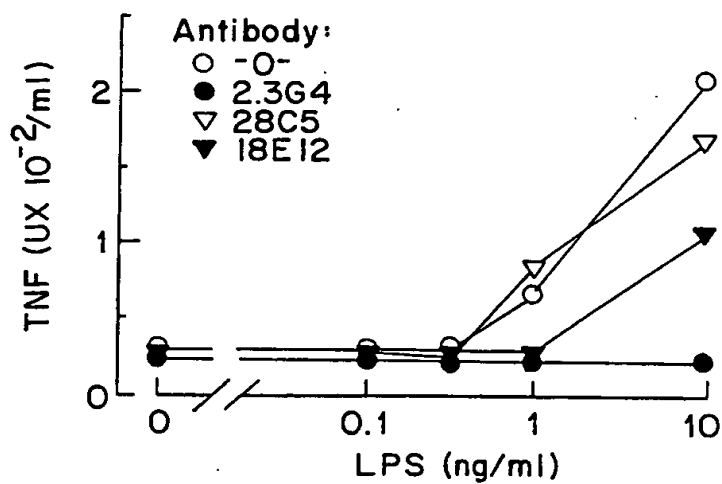


FIG. 28

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LIGHT CHAINS

3C10L	Fr.1	QSPASLAVSLGORATISC	CDR1	Fr.2	WYQQKAGQPPKSSIIY
28C5L		QSPASLAVSLGORATISC			WYQQKPGQPPKLLIIY
23G4L		QSPASLAVSLGORATISC			WYQQEPGQSPKLLIIY
18E12		QTPSSLASLGDRVITISC			WYQQPGGTVKVLIIY
3C10L	CDR2	RAANLES	Fr.3	CDR3	Fr.4
28C5L		RASNLOS			QGSYEDPWT
23G4L		VASKLES			QGSNEDPTT
18E12L		YTSRLHS			QGMNEDPYT
		GVPSRFGSGSGTDYSLTISNLEQEDFATYFC			QRGDTLPWT
					FGGGTKLEIK
3C10L	CH-1	RADAAPTVSIFPPSSEQLTSGGASVWCFLNNFYPKDINVKKWKIDGSE			
28C5L		RADAAPLVSIFPPSSEQLTSGGASVWCFLNNFYPKDINVKKWKIDVSE			
18E12L		RADAAPTVSIFPPSSEQLTSGGASVWCFLNNFYPKDINVKKWKIDGSE			
3C10L	CH-1	STYSMSSTLKDEYERHNSYTCEATRKTLTLTSTSPIVKSFN			
28C5L		STYSMSSTLTTLKDEYERHNSYTCEATHKTSTSPIVKSFN			
18E12L		STYSMSSTLTTLKDEYERHNSYTCEATHKTSTSPIVKSFN			

FIG. 29

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HEAVY CHAINS

3C10H 28C5H 18E12H	FR.1 LVKPGGSLKLSVCVSGFTFS LQSGGPGLVKPSQSLSLTCTVTGYSIT LESGPGLVAPSQSLSLTCTVSGFSLT	CDR1 SYAMS SDSAWN NYDIS	FR.3 WVRQTPEKRL EHWVA WIRQFPGNRLEWVG WIRQPPGKGLEWLG	CDR2 SISGGTTYYPDNVKG YISYSGSTSYNPSLKS VIWTSGGTNYNSAFMS
3C10H 28C5H 18E12H	FR.3 RFTISRDNARNILYLQMSSLRSEDITAMYYCAR RISITRDTSKNQFFLQLNSVTITEDTATYYCVR RLSITKDNSESQVFLKMWGLQTDGTYCYCVR	CDR3 GYDDYHY GLRFAY GDGNFYLYNFDY	FR.4 WGQGTTLTVSS WGQGTTLTVSA WGQGTTLTVSS	
3C10H 28C5H 18E12H	CH-1 AKTTPPSVYPLAPGCGDITGSSVTLGCLVKGYFPEPVTVTWNSGSLSSSVHITFPALLQSGLYTMSSSVTVPPSS AKTTPPSVYPLPPGSAQTNSMVTLGCLVKGYFPEPVTVTWNSGSLSSGVHITFPVAVLQSDLYTLSSSVTVPPSS AKTTPPSVYPLAPGSAQTNSMVTLGCLVKGYFPEPVTVTWNSGSLSSGVHITFPVAVLQSDLYTLSSSVTVPPSS			
3C10H 28C5H 18E12H	TWPSQIVTICSVAHPASSTTVDDKKLEPSGPISTINPCTSSRA TWPSETVTCNVVAHPASSTKVDDKKI TWPSETVTCNVVAHPASSTKVDDKKI			

FIG. 30

INTERNATIONAL SEARCH REPORT

International application No.
PCT/US94/05898

A. CLASSIFICATION OF SUBJECT MATTER

IPC(5) : Please See Extra Sheet.

US CL : Please See Extra Sheet.

According to International Patent Classification (IPC) or to both national classification and IPC

B. FIELDS SEARCHED

Minimum documentation searched (classification system followed by classification symbols)

U.S. : 435/7.21, 240.2, 240.27, 252.3, 320.1; 530/387.3, 388.22, 388.15; 536/23.53; 424/143.1

Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched

Electronic data base consulted during the international search (name of data base and, where practicable, search terms used)
APS, CAS ONLINE, BIOSIS,

C. DOCUMENTS CONSIDERED TO BE RELEVANT

Category*	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
Y	WO, A, 86/01533 (NEUBERGER ET AL.) 13 MARCH 1986, SEE ENTIRE DOCUMENT.	9, 11-13, 15-23, 30, 32-35, 37-45
Y	WO, A, 90/07861 (QUEEN ET AL.) 26 JULY 1990, SEE ENTIRE DOCUMENT.	10, 14-23, 31, 36-45
Y	METHODS IN ENZYMOLOGY, VOLUME 178, ISSUED 1989, BETTER ET AL., "EXPRESSION OF ENGINEERED ANTIBODIES AND ANTIBODY FRAGMENTS IN MICROORGANISMS", PAGES 476-496, SEE ENTIRE DOCUMENT.	5, 6, 9, 12, 13, 11-22, 26, 27, 30, 32-44
Y	BIOTECHNOLOGY, VOLUME 8, ISSUED JULY 1990, PUTLITZ ET AL., "ANTIBODY PRODUCTION IN BACULOVIRUS-INFECTED INSECT CELLS", PAGES 651-654, SEE ENTIRE DOCUMENT.	23, 45

☒ Further documents are listed in the continuation of Box C. ☐ See patent family annex.

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A document defining the general state of the art which is not considered to be of particular relevance	*T* later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention
E earlier document published on or after the international filing date	*X* document of particular relevance; the claimed invention cannot be considered novel or cannot be considered to involve an inventive step when the document is taken alone
L document which may throw doubts on priority claim(s) or which is cited to establish the publication date of another citation or other special reason (as specified)	*Y* document of particular relevance; the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combination being obvious to a person skilled in the art
O document referring to an oral disclosure, use, exhibition or other means	*Z* document member of the same patent family
P document published prior to the international filing date but later than the priority date claimed	

Date of the actual completion of the international search
26 AUGUST 1994

Date of mailing of the international search report

SEP 08 1994

Name and mailing address of the ISA/US
Commissioner of Patents and Trademarks
Box PCT
Washington, D.C. 20231

Authorized officer

PAULA HUTZELL

Facsimile No. (703) 305-3230

Telephone No. (703) 308-0196

INTERNATIONAL SEARCH REPORT

International application No.
PCT/US94/05898

C (Continuation). DOCUMENTS CONSIDERED TO BE RELEVANT		
Category*	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
X --- Y	PROCEEDINGS OF THE NATIONAL ACADEMY OF SCIENCES USA, VOLUME 90, ISSUED APRIL 1993, PUGIN ET AL., "LIPOPOLYSACCHARIDE ACTIVATION OF HUMAN ENDOTHELIAL AND EPITHELIAL CELLS IS MEDIATED BY LIPOPOLYSACCHARIDE-BINDING PROTEIN AND SOLUBLE CD14", PAGES 2744-2748, SEE PAGES 2745 AND 2747.	1-4, 7, 24, 25, 28, 46-50,52 ----- 5, 6, 8-23, 26, 27, 29-45, 51
X --- Y	THE JOURNAL OF CLINICAL INVESTIGATION, VOLUME 90, ISSUED DECEMBER 1992, MARTIN ET AL., "LIPOPOLYSACCHARIDE BINDING PROTEIN ENHANCES THE RESPONSIVENESS OF ALVEOLAR MACROPHAGES TO BACTERIAL LIPOPOLYSACCHARIDE. IMPLICATIONS OF CYTOKINE PRODUCTION IN NORMAL AND INJURED LUNGS", PAGES 2209-2219, SEE PAGE 2210.	1-4, 7, 46 ----- 5, 6, 8-45, 47-52
X --- Y	JOURNAL OF EXPERIMENTAL MEDICINE, VOLUME 158, ISSUED JULY 1983, VAN VOORHIS ET AL., "SPECIFIC ANTIMONONUCLEAR PHAGOCYTE MONOCLONAL ANTIBODIES", PAGES 126-145, SEE PAGES 129-130.	1, 2, 5-7, 46, 47, 49, 50 ----- 3, 4, 8-45, 48, 51, 52
X --- Y	EXPERIENTIA, VOLUME 48, ISSUED 1992, GALLAY ET AL., "COMPETITION BETWEEN LPS-BINDING PROTEIN (LBP) AND ANTI-LPS ANTIBODY IN LPS-INDUCED TNF SECRETION OF HUMAN MONOCYTES (MO), PAGE A66, ABSTRACT NO. 384, SEE ENTIRE ABSTRACT.	1, 2, 7, 46 ----- 3-6, 8-45, 47-52
Y	IMMUNOLOGY TODAY, VOLUME 4, NUMBER 3, ISSUED 1983, KOZBOR ET AL., "THE PRODUCTION OF MONOCLONAL ANTIBODIES FROM HUMAN LYMPHOCYTES", PAGES 72-79, SEE ENTIRE DOCUMENT.	8, 29

INTERNATIONAL SEARCH REPORT

International application No.
PCT/US94/05898

Box I Observations where certain claims were found unsearchable (Continuation of item 1 of first sheet)

This international report has not been established in respect of certain claims under Article 17(2)(a) for the following reasons:

1. ☐ Claims Nos.:
because they relate to subject matter not required to be searched by this Authority, namely:
2. ☐ Claims Nos.:
because they relate to parts of the international application that do not comply with the prescribed requirements to such an extent that no meaningful international search can be carried out, specifically:
3. ☐ Claims Nos.:
because they are dependent claims and are not drafted in accordance with the second and third sentences of Rule 6.4(a).

Box II Observations where unity of invention is lacking (Continuation of item 2 of first sheet)

This International Searching Authority found multiple inventions in this international application, as follows:

Please See Extra Sheet.

1. ☐ As all required additional search fees were timely paid by the applicant, this international search report covers all searchable claims.
2. ☐ As all searchable claims could be searched without effort justifying an additional fee, this Authority did not invite payment of any additional fee.
3. ☐ As only some of the required additional search fees were timely paid by the applicant, this international search report covers only those claims for which fees were paid, specifically claims Nos.:
4. ☒ No required additional search fees were timely paid by the applicant. Consequently, this international search report is restricted to the invention first mentioned in the claims; it is covered by claims Nos.:
1-52

Remark on Protest

- ☐ The additional search fees were accompanied by the applicant's protest.
☐ No protest accompanied the payment of additional search fees.

INTERNATIONAL SEARCH REPORT

International application No.
PCT/US94/05898

A. CLASSIFICATION OF SUBJECT MATTER:
IPC (5):

C07K 15/28; C07H 15/12; C12N 15/00, 1/20, 5/20, 5/10; A61K 39/395; G01N 33/53

A. CLASSIFICATION OF SUBJECT MATTER:
US CL :

435/7.21, 240.2, 240.27, 252.3, 320.1; 530/387.3, 388.22, 388.15; 536/23.53; 424/143.1

BOX II. OBSERVATIONS WHERE UNITY OF INVENTION WAS LACKING

This ISA found multiple inventions as follows:

THIS INTERNATIONAL AUTHORITY HAS FOUND 2 INVENTIONS CLAIMED IN THE INTERNATIONAL APPLICATION COVERED BY THE CLAIMS INDICATED BELOW:

- I. CLAIMS 1-52 DRAWN TO HYBRIDOMAS, ANTIBODIES, NUCLEIC ACIDS, EXPRESSION VECTORS, HOST CELLS CONTAINING AN EXPRESSION VECTOR, PHARMACEUTICAL COMPOSITIONS, AND METHODS OF DETECTING CD14, CLASSIFIED IN CLASSES 530, 536 AND 435, SUBCLASSES 387.3+, 25.53 AND 7.21 AND 240.27, RESPECTIVELY .
- II. CLAIMS 53-65 DRAWN TO METHODS OF INHIBITING BINDING OF LPS TO CD14, CLASSIFIED IN CLASS 424, SUBCLASS 143.1.

AND IT CONSIDERS THAT THE INTERNATIONAL APPLICATION DOES NOT COMPLY WITH THE REQUIREMENTS OF UNITY OF INVENTION (RULES 13.1, 13.2 AND 13.3) FOR THE REASONS LISTED BELOW:

THE DETECTING METHOD OF GROUP I AND THE THERAPEUTIC INHIBITING METHOD OF GROUP II USE DIFFERENT REAGENTS AND METHOD STEPS AND THEY ACHIEVE DIFFERING OBJECTIVES. THERE ARE THUS NO ASPECTS WHICH PROVIDE THESE TWO GROUPS WITH A SPECIAL TECHNICAL FEATURE WITHIN THE MEANING OF PCT RULE 13.2 SO AS TO FORM A SINGLE GENERAL INVENTIVE GROUP.